-Virus-host interplay-

Immediate virus recognition by RIG-I and PKR

and viral counterstrategies

Inaugural-Dissertation
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Summary

Viruses are a constant threat to mankind causing diseases ranging from mild symptoms to fatal outcome.

A rapid and efficient antiviral response is therefore crucial for the survival of the host. RIG-I-like receptors (RLR) and other immune receptors, like protein kinase R (PKR), specifically detect viral RNA species in the host cytoplasm. The sensing of virus infection triggers intracellular defense mechanisms resulting in viral alertness in the infected and surrounding cells, and forms the link to the adaptive immune system. Viruses, in turn, have evolved sophisticated countermeasures to dampen the antiviral response. The molecular mechanisms involved range from a broad shut-off of the host cell metabolism to a selective interference with key components of the immune system.

For a better understanding, which viral RNA structures are detected by immune receptors like RIG-I and PKR and what kind of viral antagonists lead to their inhibition, it is crucial to be able to determine their activation status. Hence, limited protease digestion and native polyacrylamide gel electrophoresis (PAGE) were established to directly monitor RIG-I and PKR conformational switching and oligomerization upon activation, respectively.

Various studies helped to identify RIG-I stimulating RNA features in vitro, but the first viral structure triggering an antiviral interferon response in the natural context of virus infection remained to be resolved. We identified 5’triphosphorylated (5’ppp) panhandle structures packaged into nucleocapsids as physiological RIG-I agonists. Independent of virus transcription and replication, the incoming encapsidated genomes of bunyaviruses (La Crosse virus; LACV and Rift Valley fever virus; RVFV) and orthomyxovirus (influenza A virus; FLUAV) were able to stimulate RIG-I activation and an antiviral signaling cascade. Surprisingly, antiviral activity of RIG-I against FLUAV was already promoted by binding to the 5’ppp panhandle and was independent of RIG-I downstream signaling ability. In addition to RIG-I, we also identified PKR as an immune sensor of incoming nucleocapsids. PKR thereby interacts with the intergenic region (IGR) of viral genome segments using ambisense coding strategy. Association of PKR with the IGR of incoming RVFV (Bunyaviridae) and arenavirus nucleocapsids promotes PKR phosphorylation and conformational switching and hence full PKR activation.

To antagonize immediate recognition by RIG-I and PKR, viruses need to adapt. RIG-I activation by FLUAV nucleocapsids was altered by an adaptive mutation of the polymerase complex subunit PB2. Mammalian adaptation mutation PB2 E627K stabilizes the polymerase complex association with the nucleocapsid thereby preventing RIG-I recognition of the 5’ppp panhandle. Additionally, Lassa virus (Arenaviridae) nucleoprotein interacts with PKR and promotes its degradation via the proteasomal pathway.

Therefore, we identified entry of viral nucleocapsids as the first time-point of immune recognition in the natural context of virus infection and give further insights how viruses have evolved to counteract immediate recognition.
Zusammenfassung


Um einer unmittelbaren Detektion durch RIG-I und PKR zu entgehen, müssen sich Viren anpassen. So wird die RIG-I Aktivierung durch FLUAV Nukleokapside durch eine adaptive Mutation der PB2-Polymeraseuntereinheit verändert. Die Adaptionmutation PB2 E627K stabilisiert die Interaktion des FLUAV Polymerasekomplexes mit dem Nukleokapsid und verhindert dadurch die RIG-I vermittelte
Detektion. Zudem interagiert das Lassa Virus (**Arenaviridae**-Nukleoprotein mit PKR und induziert dessen Abbau über das Proteasom.

Somit konnte das Eintreten der viralen Nukleokapside in die Wirtszelle als erster Zeitpunkt der Immunerkennung im natürlichen Kontext der Virusinfektion nachgewiesen werden. Des Weiteren ergeben sich aus dieser Arbeit Einblicke, wie Viren sich entwickelt haben um dieser unmittelbaren Immundetektion zu entkommen.
### Abbreviations

<table>
<thead>
<tr>
<th>A...</th>
<th>ActD</th>
<th>actinomycin D</th>
</tr>
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<tbody>
<tr>
<td>Asp</td>
<td></td>
<td>aspartic acid</td>
</tr>
<tr>
<td>B...</td>
<td>BDV</td>
<td>Borna disease virus (<em>Boruviridae</em>)</td>
</tr>
<tr>
<td>C...</td>
<td>CARD</td>
<td>caspase activation and recruitment domains</td>
</tr>
<tr>
<td>CHX</td>
<td></td>
<td>cycloheximide</td>
</tr>
<tr>
<td>CL13</td>
<td></td>
<td>RVFV Cl 13</td>
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<tr>
<td>Co-IP</td>
<td></td>
<td>co-immunoprecipitation</td>
</tr>
<tr>
<td>CTD</td>
<td></td>
<td>carboxy-terminal domain</td>
</tr>
<tr>
<td>CTRL</td>
<td></td>
<td>control</td>
</tr>
<tr>
<td>D...</td>
<td>dsRBD</td>
<td>dsRNA binding domain (PKR)</td>
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<tr>
<td></td>
<td>dsRNA</td>
<td>double-stranded RNA</td>
</tr>
<tr>
<td>D.mel-2</td>
<td><em>Drosophila melanogaster</em> Schneider 2 cells</td>
<td></td>
</tr>
<tr>
<td>E...</td>
<td>E</td>
<td>glutamic acid</td>
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<tr>
<td></td>
<td>eIF2α</td>
<td>eukaryotic initiation factor 2 alpha</td>
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<tr>
<td>F...</td>
<td>FLUAV</td>
<td>influenza A virus</td>
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<tr>
<td>G...</td>
<td>Glu</td>
<td>glutamic acid</td>
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<td></td>
<td>GP</td>
<td>glycoprotein</td>
</tr>
<tr>
<td></td>
<td>GSD</td>
<td>ground state depletion microscopy</td>
</tr>
<tr>
<td>H...</td>
<td>HIV</td>
<td>human immunodeficiency virus (<em>Retroviridae</em>)</td>
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<tr>
<td></td>
<td>His</td>
<td>histidine</td>
</tr>
<tr>
<td>I...</td>
<td>IFN</td>
<td>type I interferon (IFN-alpha/beta)</td>
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<tr>
<td></td>
<td>IFNAR</td>
<td>interferon-alpha receptor</td>
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<tr>
<td></td>
<td>IGR</td>
<td>intergenic region</td>
</tr>
<tr>
<td></td>
<td>IKKε/α/β</td>
<td>inhibitor of nuclear factor kappa-B kinase subunit ε/α/β</td>
</tr>
<tr>
<td></td>
<td>IRES</td>
<td>internal ribosomal entry site</td>
</tr>
<tr>
<td></td>
<td>IRF3/7</td>
<td>IFN regulatory factor 3/7</td>
</tr>
<tr>
<td></td>
<td>ISG</td>
<td>IFN stimulated gene</td>
</tr>
<tr>
<td></td>
<td>IVM</td>
<td>Ivermectin</td>
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<tr>
<td>J...</td>
<td>JAK</td>
<td>Janus kinase</td>
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<tr>
<td></td>
<td>JUNV</td>
<td>Junin virus (<em>Arenaviridae</em>)</td>
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<td>K...</td>
<td>K</td>
<td>lysin</td>
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<tr>
<td>L...</td>
<td>LACV</td>
<td>La Crosse virus (<em>Bunyaviridae</em>)</td>
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<td></td>
<td>LASV</td>
<td>Lassa virus (<em>Arenaviridae</em>)</td>
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<tr>
<td></td>
<td>LGP2</td>
<td>laboratory of genetics and physiology 2</td>
</tr>
<tr>
<td></td>
<td>LMB</td>
<td>Leptomycin B</td>
</tr>
<tr>
<td>Abbreviation</td>
<td>Description</td>
<td></td>
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<tr>
<td>--------------</td>
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<tr>
<td>MAVS</td>
<td>mitochondrial antiviral signaling protein</td>
<td></td>
</tr>
<tr>
<td>MAM</td>
<td>mitochondrial associated membrane</td>
<td></td>
</tr>
<tr>
<td>MAPK</td>
<td>mitogen activated protein kinase</td>
<td></td>
</tr>
<tr>
<td>MDA5</td>
<td>melanoma differentiation association factor 5</td>
<td></td>
</tr>
<tr>
<td>N…</td>
<td>nucleoprotein (Bunyaviridae)</td>
<td></td>
</tr>
<tr>
<td>NF-κB</td>
<td>nuclear factor κ-light-chain-enhancer of activated B cells</td>
<td></td>
</tr>
<tr>
<td>NP</td>
<td>nucleoprotein ( Arenaviridae and Orthomyxviridae)</td>
<td></td>
</tr>
<tr>
<td>NS1</td>
<td>non-structural protein 1 (FLUAV)</td>
<td></td>
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<tr>
<td>NSs</td>
<td>non-structural protein encoded on the S segment (RVFV)</td>
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<tr>
<td>NSV</td>
<td>negative-strand RNA virus</td>
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<tr>
<td>P…</td>
<td>triphosphate</td>
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<tr>
<td>PA</td>
<td>polymerase acidic protein</td>
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<tr>
<td>PACT</td>
<td>protein activator of PKR</td>
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<td>PAGE</td>
<td>polyacrylamide gel electrophoresis</td>
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<td>PB1/2</td>
<td>polymerase basic protein 1/2</td>
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<td>PCR</td>
<td>polymerase chain reaction</td>
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<tr>
<td>PHV</td>
<td>Prospect Hill virus (Bunyaviridae)</td>
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<tr>
<td>PKR</td>
<td>protein kinase R</td>
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<tr>
<td>Q…</td>
<td>quantitative reverse transcription polymerase chain reaction</td>
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<tr>
<td>R…</td>
<td>retinoic acid inducible gene I</td>
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<tr>
<td>RLR</td>
<td>RIG-I like receptor</td>
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<tr>
<td>RNA</td>
<td>ribonucleic acid</td>
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<tr>
<td>RVFV</td>
<td>Rift Valley fever virus (Bunyaviridae)</td>
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<tr>
<td>S…</td>
<td>shrimp alkaline phosphatase</td>
<td></td>
</tr>
<tr>
<td>siRNA</td>
<td>small interfering RNAs</td>
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</tr>
<tr>
<td>STAT</td>
<td>signal transducer and activator of transcription</td>
<td></td>
</tr>
<tr>
<td>T…</td>
<td>transactivation RNA (HIV)</td>
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</tr>
<tr>
<td>TBK1</td>
<td>TANK-binding kinase 1</td>
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<td>TCRV</td>
<td>tacaribe virus ( Arenaviridae)</td>
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<td>TRAF 3/6</td>
<td>TNF receptor-associated factor 3/6</td>
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<td>TRIM25</td>
<td>tripartite motif containing 25</td>
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<tr>
<td>U…</td>
<td>56 kDa U2AF65-associated protein</td>
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<tr>
<td>URH49</td>
<td>UAP56-related helicase, 49 kDa</td>
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<tr>
<td>W…</td>
<td>wildtype</td>
<td></td>
</tr>
<tr>
<td>Z…</td>
<td>matrixprotein (Lassa virus)</td>
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1. Introduction

1.1. Induction of antiviral defense mechanisms by RIG-I like receptors

Viruses are a constant threat to mankind. Thereby, most of the known emerging and highly pathogenic viruses are RNA-genome based. They give rise to epidemic and rarely pandemic diseases, as often reported for influenza viruses [30]; or cause hemorrhagic fever, like members of the families Arenaviridae (Lassa fever virus and Junin virus) [35, 147] and Bunyaviridae (Rift Valley fever virus and Crimean Congo haemorrhagic fever virus) [6, 51]. Thus, to rapidly control virus infection an immediate recognition of the viral intruder is required.

1.1.1. RIG-I like receptors

The germ-line encoded RLR (RIG-I (retinoic acid inducible gene I) like receptors) represent a group of cytoplasmic sensor proteins able to detect RNA virus infection. RLR belong to the DExD/H-box helicase family (refers to Asp-Glu-x-Asp/His, where x can be any amino acid) within the helicase superfamily 2. RIG-I, the closely related MDA5 (melanoma differentiation association factor 5) and LGP2 (laboratory of genetics and physiology 2) form the family of RLR. RIG-I and MDA5 complement each other by responding to diverse virus families and initiate antiviral type I interferon (IFN-alpha/beta) responses [144]. The role of LGP2 in cytosolic virus sensing is less well characterized and currently controversially discussed. However, it is widely accepted that LGP2 represents rather a modulator of RIG-I and MDA5 activity then being involved in immune sensing or signaling [107, 109, 151].

1.1.2. RIG-I like receptor agonists

Studies have demonstrated that RIG-I and MDA5 recognize mainly distinct virus families. RIG-I is required to initiate an antiviral response against Filoviridae (Ebola and Marburg virus), Paramyxoviridae (New Castle Disease virus and respiratory syncytial virus), Rhabdoviridae (vesicular stomatitis virus), Orthomyxoviridae (Influenza virus), Bunyaviridae (Rift Valley Fever virus) and Flaviviridae (Japanese encephalitis virus). On the other hand, MDA5 responds to representatives of the Picornaviridae (Encephalomyocarditis virus). Together, RIG-I and MDA5 react to Reoviridae and representatives of the Flaviviridae group (West Nile virus and Dengue virus) [112, 145]. An essential feature of the RLR is specificity to distinguish between host cell (self) and viral (non-self) patterns. Thereby, MDA5 is known to sense longer dsRNA (double-stranded RNA) molecules up to two kilo bases [58, 92], ideally with higher order RNA structures [92]. RIG-I mainly responds to 5’ ppp blunt-ended dsRNA of a minimal length of 10 base pairs [46, 60, 91, 111, 113]. Also dsRNA stretches bearing a 5’ diphosphate were recently identified to stimulate a RIG-I dependent response [33]. Long dsRNA molecules of more than 200 base pairs (irrespective of the 5’ ends) [9], 3’-monophosphorylated cleavage products of RNase L [73], and polyuridine or polyriboadenine stretches were also described as
RIG-I agonists [103, 108, 112]. Importantly, these RNA structures and modalities are absent in the cytoplasm of non-infected cells, allowing the discrimination between self and non-self.

1.1.3 Structure-based RIG-I activation
All RLR share a similar structure (manuscript 4.5, Fig. 2). Thereby, RIG-I is composed of a CTD (carboxy-terminal domain) and a conserved RNA helicase core (formed by Hel1 and Hel2 with the insertion domain Hel2i). Both domains are required for agonist recognition in which the CTD has high affinity for the 5’ppp and the helicase domain recognizes dsRNA [128]. The helicase domain also promotes ATP hydrolysis. The CTD is connected to Hel2 via the pincer domain (also named bridging domain), which transmits the information of ligand recognition to the helicase core [57, 65, 71, 98]. Additionally, RIG-I possesses two amino-terminal CARDs (caspase activation and caspase recruitment domains) for signal transduction [64, 72, 144]. MDA5 has a similar structure whereas LGP2 lacks CARDs for downstream signaling [139, 144].

In resting state, RIG-I is present as an extended monomer in an open auto-repressed conformation [62, 71]. Thereby, the CARDs contact each other and CARD2 interacts with Hel2i, which sterically impedes ligand and coactivator engagement and initiation of downstream signaling [65, 97]. The CTD is, however, flexibly exposed and surveys the environment for 5’ppp-dsRNA. CTD binding to the 5’ppp brings the dsRNA in close proximity to the helicase domain [62]. The helicase core cooperatively binds the phosphate backbone of the dsRNA and ATP and the associated conformational switch promotes the release of CARDs [65]. CARDs are then freely accessible for polyubiquitinylation by the E3 ligase TRIM25 [29] or unanchored polyubiquitin chains [149]. Polyubiquitinylation prevents reassociation of the CARDs with the helicase domain to return into the auto-repressed state [62]. Furthermore, ubiquitinylation allows association of RIG-I molecules into a tetrameric complex [56, 89] leading to the fully signaling competent RIG-I complex.

1.1.4 RIG-I mediated type I interferon response
Activated RIG-I complexes recruit the adaptor protein MAVS (mitochondrial antiviral signaling) via CARD-CARD interaction [140]. Thereby, a transient interaction is sufficient to promote self-perpetuating MAVS filament association and thus, leading to a large-scale amplification of the antiviral signaling cascade [47]. Although the majority of MAVS is present on the mitochondria, it is also localized to peroxisomes [23]. Both organelles act sequentially as signaling platforms. Thereby, peroxisomes induce a rapid and transient IFN-independent expression of ISGs whereas mitochondrial MAVS initiates a sustained IFN-dependent response with delayed kinetics [23]. MAM (mitochondrial associated endoplasmatic reticulum membrane) directs relocalization of MAVS between the organelles, thus coordinates signaling between peroxisomes and mitochondria [45]. MAVS filament formation initiates the recruitment of several molecules to assemble a signaling platform [47, 55, 142]. Interaction of MAVS with TRAF3 and TRAF6 (TNF receptor-associated factor 3 and 6, respectively)
promotes activation of TBK1 (TANK-binding kinase 1) and IKKε (inhibitor of nuclear factor kappa-B kinase subunit ε) responsible for the phosphorylation of IRF3 and IRF7 (IFN regulatory factor 3 and 7, respectively), and IKKα and IKKβ promote NF-κB (nuclear factor κ-light-chain-enhancer of activated B cells) activation. Phosphorylated IRF3/7 forms dimers and translocates, like NF-κB, into the nucleus to initiate expression of IFN, proinflammatory cytokines and RIG-I and MDA5, providing a positive feedback mechanism for the amplification of the antiviral response [32, 103]. IFN binding to IFNAR (interferon-alpha receptor) stimulates the JAK (janus kinase)-STAT (signal transducer and activator of transcription) signaling pathway leading to selective transcriptional activation of numerous IFN-stimulated genes (ISGs) [44, 52]. ISGs represent effector proteins of the IFN response, which directly act against multiple steps of virus replication to establish an antiviral state in infected and neighboring cells [87, 116]. Furthermore, costimulatory molecules, cytokines and chemokines favor the initiation of adaptive immune responses [70, 115, 117]. The innate immune system thus controls virus infection at early phases and permits a subsequent specific adaptive immune response to clear infection and establish an immunogenic memory [41].

1.1.5. Protein kinase R
One well studied effector protein of the innate immune response is PKR (protein kinase R). PKR is expressed at a low constitutive level in the cytoplasm, but can casually be detected in the nucleus. To stimulate PKR activation there is a minimal length requirement of dsRNA of approximately 16 base pairs, but also specific structural modalities, like RNA bulges and loops, and nucleotide modifications, like a 5’ppp, support PKR stimulation [7, 25, 84]. PKR thereby contacts the dsRNA in a sequence-independent manner [7]. However, during stress PACT (protein activator of PKR) can activate PKR in absence of a stimulating dsRNA ligand [88]. Concerning PKR structure, the protein is composed of two amino-terminal dsRBD (dsRNA binding domains), a carboxy-terminal serine/threonine kinase domain, whereby both domains are linked by a flexible linker [126]. In absence of a stimulus, PKR is present as a monomer with an extended, open conformation and the dsRBDs are impeding kinase domain activity. Agonist recognition induces structural rearrangements to a closed conformation allowing dimerization and auto-phosphorylation [16-18, 67, 126]. Once activated, PKR is a multifunctional protein involved in the regulation of cap-dependent translation, it participates in the formation of virus induced stress granules required for IFN signaling and controls NF-κB, p38 MAPK and insulin pathways, among others [18, 22, 85, 148].

1.2. Viral evasion strategies to prevent immune recognition
Induction of the IFN response acts as a major barrier to virus infection, which must be circumvented to enable a productive infectious cycle. Thus, viruses have evolved multiple mechanisms to target key signaling molecules of the RLR pathway to prevent establishment of an antiviral state. A common strategy of RNA viruses comprises the modification and concealing of the viral genomes and
replicative intermediates. One mechanism presents packaging of the viral RNA with multiple copies of viral nucleoprotein and the polymerase into nucleocapsids [106]. Furthermore, some viruses possess the ability to enzymatically remove their 5`ppp in order to avoid RIG-I detection [36]. Other viruses hide their genomes by replicating in the nucleus or in inaccessible cellular compartments [152]. In course of the viral replication cycle, accumulation of erroneous, misencapsidated replication products might serve as RLR agonists. Viruses therefore mask [41, 42, 95], degrade these erroneous side products [38, 41, 63, 93] or use cellular helicases to unwind dsRNA replication intermediates [136, 137].

Overall, viruses have evolved sophisticated strategies to prevent RLR recognition and the subsequent IFN response. Further knowledge of viral evasion strategies is, however, required to comprehend the complexity of virus-host interactions to gain insights into RLR signaling and also virus pathogenesis itself.
1.3. Objectives of the underlying thesis

Despite the discoveries about stimulating RIG-I structures, the question of the physiological RIG-I agonist in the authentic infection context remained largely unresolved. Long dsRNA is not produced in a detectable amount during the course of negative-strand RNA virus (NSV) infection [131]. However, NSV possess complementary sequences at the termini of their genomes leading to the formation of the “panhandle” conformation with a short dsRNA stretch. Furthermore, because of initiation of RNA synthesis via a single nucleoside triphosphate, all NSV were believed to possess a 5′ppp at the termini of their genomes [36]. Therefore, the 5′ppp panhandle structure represents all hallmarks of a potential RIG-I agonist. Nevertheless, the genomes of NSV do not exist as free RNA, but are instead hidden in a complex with multiple copies of viral nucleoprotein and the polymerase, the so-called nucleocapsid. It has remained unclear, whether RIG-I can also recognize RNAs packaged into nucleocapsids, the main viral RNA state in the infected host cell.

First aim of this study was hence the identification of the physiological RIG-I agonist during virus infection. Thereby, we aimed to pin down the first viral structure to be detected by RIG-I and able to stimulate the host innate immune system. We focused at first on stimulating RNA structures of the cytoplasmic bunyaviruses and extended then our study towards influenza viruses, which are only briefly accessible to RIG-I during their cytoplasmic transit to the nucleus for genome replication.

Besides RIG-I, we wondered whether other immune receptors would promote an immediate antiviral response. PKR is able to respond to dsRNA, but also specific structural modalities, like bulges and loops [25]. Interestingly, intergenic regions of viral genomes using ambisense coding strategies resemble loop structures with a dsRNA stem. Thus, we hypothesized that PKR could be able to induce immediate antiviral defense mechanisms by recognizing intergenic regions of incoming viral genomes.

To validate this hypothesis was the second aim of this thesis.

On the contrary, to counteract immediate recognition viruses have evolved sophisticated methods. Viral immune evasion strategies thereby range from an unspecific shut-off of the host cell metabolism to a selective interference with key components of the IFN system. In our research, we intended to assess whether NSVs have evolved so far unknown mechanisms to specifically evade immediate immune recognition.

Overall, we intend to provide further insights in immediate virus detection in the natural context of infection and how viruses, on the other hand, have adapted to evade immune recognition to allow efficient virus replication and spread.
2. Results

2.1. Monitoring the activation status of RIG-I and PKR

For a better understanding, which viral RNA structures are detected by immune receptors, like RIG-I and PKR, and what kind of viral antagonists lead to their inhibition, it is crucial to be able to determine their activation status. So far, techniques to measure downstream events like antiviral gene expression are commonly used, but since viruses interfere with RLR and IFN signaling, results might not reflect the direct activation status of RIG-I or PKR. Here, we present limited protease digestion and native polyacrylamide gel electrophoresis (PAGE) as sensitive and direct measurements of two distinct markers of RIG-I and PKR activation (manuscript 4.1, [133]). Upon agonist recognition RIG-I and PKR undergo conformational switching [2, 107, 121] allowing formation of oligomeric complexes [22, 56, 107]. Sensitive validation of the RIG-I and PKR activation status helps to gain further insights in physiological agonists and how viruses interfere with RIG-I and PKR activation. In absence of a specific agonist, RIG-I has an open auto-repressed conformation [62, 71]. Agonist recognition induces conformational switching of RIG-I and formation of a much more ordered RIG-I molecule [62, 72]. PKR is also present as a monomer with an extended open conformation in the absence of a stimulus. Binding to a specific agonist leads to structural rearrangements to a closed conformation allowing dimerization and auto-phosphorylation [17, 67, 126]. Conformational rearrangements lead to alterations in protease sensitivity, which can be determined by limited protease digestion. This technique, to monitor RIG-I and PKR conformational switching, was previously described by M. Gale Jr. [107] and T. Fujita [121], and J.L. Cole [2], respectively. To monitor RIG-I and PKR conformational switching upon virus infection, human A549 cells were either mock infected or infected with Rift Valley fever virus clone 13 (Cl 13). Cl 13 lacks a functional IFN antagonist NSs and is hence a strong activator of RIG-I and PKR [36, 37, 50]. Trypsin digestion of mock infected cell lysates results in a rapid degradation of RIG-I whereas Cl 13 infection leads to the generation of a 30 kDa protease resistant RIG-I fragment (manuscript 4.1, Fig. 1A). Also PKR shows partial resistance to trypsin digestion in Cl 13 infected samples, which coincides with its phosphorylation at threonine 466, a widely accepted marker of PKR activation (manuscript 4.1, Fig. 1A).

RIG-I and PKR conformational rearrangements form the prerequisite to associate multiple monomers into oligomeric complexes. Accumulation of RIG-I oligomers allows enhanced recruitment of downstream signaling molecules to form a platform for antiviral signal transduction [72]. In case of PKR, dimerization supports auto-phosphorylation required for its full activation [17, 22]. The formation of oligomeric complexes was assayed by native PAGE. Proteins and protein complexes are thereby not separated by their charge to mass ratio, but rather by their size [138].

By applying native PAGE, formation of RIG-I and PKR oligomeric complexes in Cl 13 infected cell lysates could be demonstrated whereas these proteins remained as monomers in mock infected
samples (manuscript 4.1, Fig. 2, upper and middle panel). As an additional control for the functionality of the oligomerization assay the IFN transcription factor IRF3 was included. Validation of IRF3 dimerization is widely performed to monitor activation of the IFN system [54]. Indeed, IRF3 dimerization coincides with RIG-I and PKR oligomerization in C13 infected samples (manuscript 4.1, Fig. 2, lower panel).

Taken together, these results demonstrate limited trypsin digestion and native PAGE as useful and sensitive tools to directly monitor two distinct markers of RIG-I and PKR activation.

2.2. Characterization of the physiological RIG-I agonist

2.2.1. Incoming bunyavirus nucleocapsids as RIG-I agonists

Most studies on the identification of RIG-I agonist were performed by allowing full viral replication or by transfecting naked viral or synthetic RNA. These results helped to gain insights into RNA features required for RIG-I activation, like dsRNA and a 5’ppp, but did not answer the question about the RIG-I activating structure in the natural virus infection. Therefore, the first objective of this thesis was the identification of the earliest infection step exposing a viral RNA pattern able to activate RIG-I and stimulate an antiviral IFN response in an authentic virus infection (manuscript 4.2, [132]).

The infection cycle of NSV starts with particle attachment to the host cell, followed by entry of the viral genome into the cytoplasm, mRNA (messenger RNA) synthesis during primary transcription via the viral polymerase, genome replication via a positive-sense template, assembly and finally release of progeny viruses. It has been demonstrated that full-length and defective RNAs produced during viral replication serve as RIG-I stimulators [5, 101]. However, whether these RNA structures represent naked erroneous side products of viral replication was not addressed. During virus infection, the viral genomic and intermediate RNA products never exist as free RNA, but are instead hidden within the nucleocapsid [131]. So far, it has remained unresolved whether RIG-I can also recognize RNAs packaged into nucleocapsids, the main viral RNA state in the infected host cell.

Nucleocapsids of NSVs represent hallmarks of a potential RIG-I agonist. Terminal complementary sequences of the viral genomes allow hybridization into a short dsRNA stretch, classified as the “panhandle” conformation. Moreover, the 5’ppp is generated during initiation of RNA synthesis via a single nucleoside triphosphate [110]. To study whether the incoming 5’ppp nucleocapsids would suffice to trigger an IFN response the peculiar characteristics of the bunyaviruses were exploited. Bunyavirus transcription is dependent on on-going translation [94]. Hence, treatment with the translation inhibitor CHX (cycloheximide) aborts primary transcription and allows validation of incoming nucleocapsids as IFN stimulators. Infections were performed with two representatives of the Bunyaviridae, RVFV (Rift valley fever virus) and LACV (La Crosse virus). To test induction of an IFN response, IFN-beta and ISG56 mRNA upregulation were measured by qRT PCR (quantitative reverse transcription PCR). Results indicate that independent of primary transcription or genome replication, already the incoming LACV and RVFV nucleocapsids stimulate an IFN response.
Results

To validate the influence of the 5’ppp, PHV (Prospect Hill virus), another bunyavirus, which processes its 5’terminus to a 5’p (monophosphate) [31, 36] was included into the study. This allows a direct comparison of the antiviral response by nucleocapsids with (LACV and RVFV) or without (PHV) the 5’ppp. By performing the experiment with PHV no upregulation of IFN-beta and ISG56 was detected (manuscript 4.2, Fig. 2A). This indicates that the induction of the IFN response by nucleocapsids seems to be dependent on the presence of a 5’ppp.

To further validate induction of the IFN response by 5’ppp nucleocapsids, the activation status of the IFN transcription factor IRF3 was analyzed. Thereby, phosphorylation, dimerization and translocation of IRF3 into the nucleus represent central features of its activation [143]. Indeed, LACV and RVFV nucleocapsids promote phosphorylation, dimerization (manuscript 4.2, Fig. S3D and S4D, respectively) and translocation of IRF3 (manuscript 4.2, Fig. S3E) whereas PHV 5’p nucleocapsids do not stimulate a detectable IRF3 activation (manuscript 4.2, Fig. S3F). This data supports the previous result that triggering an antiviral IFN response requires 5’ppp nucleocapsids.

To test which RLR is responsible for the induction of the immune response, knockdown cells were generated by transfecting siRNA (small interfering RNAs) against non-specific target (CTRL), MDA5 or RIG-I. Knockdown of MDA5 did not affect the ability of incoming LACV nucleocapsids to stimulate IFN-beta and ISG56 expression (manuscript 4.2, Fig. 2B). However, absence of RIG-I resulted in an impaired IFN response. This indicates that RIG-I is required to sense nucleocapsids. Taken together, incoming 5’ppp nucleocapsids promote a RIG-I dependent antiviral IFN response.

To analyze whether RIG-I is able to interact with the incoming nucleocapsids, confocal immunofluorescence microscopy and co-IP (co-immunoprecipitation) assay were performed. Detection of the nucleoprotein N served thereby as a marker for the viral nucleocapsids. Confocal immunofluorescence microscopy revealed that RIG-I colocalized with LACV and RVFV nucleocapsids (manuscript 4.2, Fig. 4A and S4E). Incoming nucleocapsids also showed a colocalization with peroxisomes (manuscript 4.2, Fig. S4B), which are known organelles for induction of an immediate antiviral response [23]. This interaction between RIG-I and incoming nucleocapsids was further validated by co-IP. With a RIG-I specific co-IP, also LACV nucleocapsids could be precipitated (manuscript 4.2, Fig. 4B). This demonstrates that RIG-I is able to interact with incoming nucleocapsids. Whether RIG-I interaction with the incoming nucleocapsids would promote RIG-I activation was validated by the previously described assays (manuscript 4.1, [133]). Therefore, we performed limited protease digestion and native PAGE to monitor RIG-I conformational switching and oligomerization, respectively. Full viral replication and incoming LACV and RVFV nucleocapsids induced trypsin resistant RIG-I fragments as a marker of conformational switching (manuscript 4.2, Fig. 3G and S4D). However, PHV full replication cycle and PHV incoming 5’p nucleocapsids only poorly stimulated RIG-I conformational changes. Also RIG-I oligomerization increased upon full LACV and RVFV replication and remained at a constant, but clearly detectable level if replication cycle was aborted after entry of the viral nucleocapsids (manuscript 4.2, Fig. 3F
and S4D). In contrast, PHV infection and incoming 5’p nucleocapsids failed to promote accumulation of RIG-I oligomeric complexes (manuscript 4.2, Fig. 3H).

An insect cell system was additionally employed to investigate whether RIG-I activation by viral nucleocapsids is direct or mediated by one of the abundant cellular cofactors of RIG-I [61, 81, 87, 146]. Therefore, isolated nucleocapsids from purified RVFV or LACV virions were used to stimulate human RIG-I expressed from D.mel-2 (Drosophila melanogaster Schneider 2) cells. In absence of mammalian cofactors, purified nucleocapsids promote RIG-I conformational switching (manuscript 4.2, Fig. 3H). Thus, mammalian cofactors are not required for nucleocapsid-stimulated RIG-I activation. To test the requirement of the 5’ppp and dsRNA for RIG-I activation by incoming nucleocapsids, purified nucleocapsids were either treated with a ssRNA-specific RNase A, a dsRNA-specific RNase III or SAP (shrimp alkaline phosphatase). By destroying the dsRNA stretch (RNase III) or removal of the 5’ppp (SAP), the nucleocapsids lost their ability to promote RIG-I conformational switching, whereas RNase A treatment did not have an effect (manuscript 4.2, Fig. 6E). This indicates that the 5’ppp dsRNA panhandle is required for RIG-I activation. Furthermore, by employing super-resolution GSD (ground state depletion) microscopy we were able to display the pseudocircular structure of LACV nucleocapsids and that accumulations of RIG-I contact the nucleocapsids via a single contact site, most likely the 5’ppp dsRNA panhandle (manuscript 4.2, Fig. 7).

In summary, RIG-I interacts with incoming nucleocapsids comprising a 5’ppp dsRNA panhandle. This interaction leads to RIG-I conformational switching and oligomerization independent of other mammalian cofactors. Once activated, RIG-I promotes, partially via the peroxisomal signaling platform, activation of the IFN transcription factor IRF3 and the antiviral type I IFN response. This defines nucleocapsid entry as the first time-point when the host immune system encounters the viral intruder to promote an immediate antiviral response.

### 2.2.2. Incoming influenza virus nucleocapsids serve as RIG-I agonists

Influenza A viruses (FLUAV) belong to the family Orthomyxoviridae containing a segmented, single-stranded RNA genome with negative-strand polarity. Each FLUAV particle comprises eight genome segments encapsidated by the viral nucleoprotein NP and the polymerase complex subunits PB1, PB2 and PA. The 5’ and 3’ ends of each viral RNA contain partially complementary sequences forming, as described for other NSVs, a panhandle conformation with a short dsRNA [48, 90]. Since incoming 5’ppp nucleocapsids of the bunyaviruses could be identified as physiological RIG-I agonists (manuscript 4.2, [132]), we wondered if also FLUAV nucleocapsids would serve as RIG-I stimulators. However, FLUAV transcription and replication occur in the nucleus. Due to the viral replication cycle, the incoming FLUAV nucleocapsids are only shortly exposed to cytoplasmic immune receptors. Here, we addressed the question if the short cytoplasmic exposure of FLUAV
nucleocapsids during their transit to the nucleus is sufficient to activate RIG-I (manuscript 4.3, (Weber et al., in press)).

To study, whether RIG-I is able to recognize incoming FLUAV nucleocapsids, diverse inhibitors were used to restrict virus infection. CHX (cycloheximide) blocks protein synthesis and therefore viral genome replication, LMB (Leptomycin B) inhibits nuclear export of nucleocapsids, ActD (actinomycin D) is an inhibitor of viral transcription, and IVM (Ivermectin) is known to block nuclear import of viral nucleocapsids [130]. Furthermore, to ensure analysis of an immediate response infection was stopped one hour post infection. Indeed, independent of the applied inhibitor, incoming A/PR/8/34 (H1N1) nucleocapsids promote RIG-I conformational switching and oligomerization and thus its full activation (manuscript 4.3, Fig. 1A and B).

To verify an interaction between RIG-I and the incoming FLUAV nucleocapsids various approaches were applied. Super-resolution GSD microscopy revealed that RIG-I is attached to the rod-like FLUAV nucleocapsids via a single contact site (manuscript 4.3, Fig. 2A). Association of RIG-I with FLUAV nucleocapsids was further confirmed by co-IP and co-sedimentation assays. Thereby, FLUAV nucleocapsids could be coprecipitated with a RIG-I specific IP (manuscript 4.3, Fig. 2B). Likewise, by performing co-sedimentation assay, RIG-I and FLUAV nucleocapsids partially shifted together into the same fractions (manuscript 4.3, Fig. 2C). This indicates that RIG-I is indeed able to interact with incoming FLUAV nucleocapsids.

To test the influence of mammalian RIG-I cofactors the D-mel.2 in vitro system was applied. Also in absence of any mammalian factors, purified FLUAV nucleocapsids induced RIG-I conformational switching (manuscript 4.3, Fig. S2F), oligomerization (manuscript 4.3, Fig. S2G) and a RIG-I shift in a co-sedimentation assay (manuscript 4.3, Fig. S2H). Therefore, also in absence of other mammalian factors RIG-I interacts and gets activated by FLUAV nucleocapsids. RIG-I activation is, however, dependent on the presence of a 5’ppp and a dsRNA structure since the FLUAV nucleocapsids lose their RIG-I activating potential after treatment with a dsRNA-specific RNase III or the phosphatase SAP, respectively (manuscript 4.3, Fig. 2D).

To validate whether FLUAV nucleocapsids can promote an IFN response, the activation status of IRF3 was validated. IRF3 phosphorylation, one marker of its activation, was induced upon presence of FLUAV nucleocapsids (manuscript 4.3, Fig. 1D).

As a conclusion, incoming FLUAV nucleocapsids serve as natural RIG-I agonist independent of RNA synthesis or presence of mammalian cofactors, triggering a signaling cascade that culminates in the activation of the IFN transcription factor IRF3.

### 2.3 PKR as an immediate sensor of virus infection

As previous results indicate, incoming RVFV and LACV (Bunyaviridae) and FLUAV (Orthomyxoviridae) nucleocapsids serve as physiological RIG-I agonists enabling an immediate antiviral type I IFN response (manuscript 4.2, [132] and manuscript 4.3, (Weber et al., in press)).
However, whether other host immune receptors would likewise contribute to immediate virus recognition remained unresolved.

One well-characterized cytoplasmic immune sensor is PKR. With its ability to recognize dsRNA stretches or specific structural RNA modalities, like bulges and loops [18], PKR comprises the potential to recognize dsRNA replicative intermediates of NSVs [20]. However, whether PKR can be activated by incoming viral RNA structures packaged within the nucleocapsids was not addressed so far (manuscript 4.4, Weber et al., manuscript in preparation).

Upon agonist recognition, conformational rearrangements of PKR lead to partial protease resistance allowing dimerization and auto-phosphorylation (manuscript 4.1, [133]). By comparing the potential of the two closely related LACV and RVFV to promote PKR conformational switching and phosphorylation, interesting differences were observed. LACV full infectious cycle and incoming LACV nucleocapsids stimulate only weakly these two markers of PKR activation (manuscript 4.4, Fig. 1B). On the contrary, full RVFV replication and incoming nucleocapsids of RVFV lead to a robust detection of PKR conformational switching and phosphorylation. Interaction of PKR with bunyavirus nucleocapsids was validated by co-sedimentation assay and co-IP. By performing co-sedimentation assay, PKR shifted to similar fractions as RVFV nucleocapsids whereas upon stimulation with LACV nucleocapsids PKR showed a similar distribution in the gradient as upon mock infection (manuscript 4.4, Fig. 1C). Co-IPs confirmed PKR interaction with RVFV, but not LACV, nucleocapsids (manuscript 4.4, Fig. 1D). This indicates that incoming RVFV nucleocapsids can be recognized by PKR and stimulate its activation. Hence, incoming RVFV nucleocapsids serve as natural PKR agonists.

To identify the PKR stimulating structure present within RVFV, but absent in LACV nucleocapsids, the coding strategies of these two viruses were analyzed. Bunyaviruses divide their genome into three segments, which are named L (large), M (middle) and S (small) according to their size [11]. RVFV L and M segment and all segments of LACV are present in negative polarity with one transcriptional unit coding for one or more proteins. RVFV S segment, however, uses ambisense coding strategy (manuscript 4.4, Fig. 2B). Thereby, two open reading frames in opposite directions are separated by a non-coding intergenic region (IGR). RVFV IGR resembles a hairpin structure with a central dsRNA stem disordered by internal loops (manuscript 4.4, Fig. 2A). To test whether this peculiar structure is responsible for PKR activation, RVF VLP (RVF virus like particles) containing either M (no IGR) or S (with IGR) segmented nucleocapsids were generated. Due to the same 5'ppp dsRNA panhandle of M and S segments the nucleocapsids promote RIG-I conformational switching to the same extent (manuscript 4.4, Fig. 2C). However, only S segmented nucleocapsids possessing the IGR are able to stimulate PKR conformational switching. As detected by super-resolution GSD microscopy, RIG-I colocalized with both M and S segmented nucleocapsids whereas PKR only showed interaction with S segmented nucleocapsids (manuscript 4.4, Fig. 2D). Taken together, PKR activation by incoming nucleocapsids requires the presence of a structured IGR.
To further support this result, arenavirus nucleocapsids were validated based on their potential to promote PKR activation. According to antigenicity, phylogeny and geographical distribution, arenaviruses are classified into Old World and New World viruses [63]. Arenavirus particles comprise two genome segments both using ambisense coding strategy with an IGR for transcriptional control. Compared to RVFV IGR, representatives of the arenaviruses, like the Old World LASV (Lassa virus) and New World JUNV (Junin virus) and TCRV (Tacaribe virus) possess also a highly structured IGR with a central dsRNA stem (manuscript 4.4, Fig. 3A). By validating PKR activation, TCRV, JUNV and LASV nucleocapsids induce PKR phosphorylation and conformational rearrangements (manuscript 4.4, Fig. 3B, right panel). Thus, incoming arenavirus nucleocapsids serve, like RVFV nucleocapsids, as natural PKR agonists.

This data demonstrate that PKR is able to recognize IGRs exposed from the incoming nucleocapsid complex. IGR engagement promotes PKR conformational switching and phosphorylation identifying IGRs of RVFV S segment, New World arenavirus TCRV and JUNV and Old World arenavirus LASV nucleocapsids as natural PKR activators. This emphasizes PKR as an immune sensor of immediate virus infection.

2.4. Viral immune evasion strategies to prevent immediate recognition

2.4.1. Segmented negative-strand RNA viruses as strong immune stimulators

The reviews included in this cumulative doctoral thesis briefly summarize the current understanding of immune evasion strategies of segmented and non-segmented NSVs (manuscript 4.5, [135] and manuscript 4.6, [134]). However, in our research, we focused on the molecular mechanisms of NSVs with a segmented genome to evade immediate immune recognition.

Besides Orthomyxoviridae with up to eight genome segments, members of the Arenaviridae and Bunyaviridae have two and three genome segments, respectively. Viruses with a segmented genome possess complementary sequences at the termini of their genomes leading to the formation of the panhandle conformation [26, 27, 76]. As outlined above, 5`ppp dsRNA panhandle structures of incoming bunyavirus (manuscript 4.2, [132]) and influenza virus (manuscript 4.3, (Weber et al., in press)); and arenavirus nucleocapsids (manuscript 4.4, (Weber et al., manuscript in preparation)) serve as natural RIG-I and PKR agonists, respectively. Hence, NSVs with segmented genomes immediately present an increased number of potential immune receptor agonists leading to an early activation of antiviral defense mechanisms (manuscript 4.5, Fig. 3). To evade early immune recognition and the subsequent antiviral IFN response sophisticated countermeasures are thus required. Here, we address the question how influenza viruses and arenaviruses prevent immediate nucleocapsid detection by the host immune receptors RIG-I and PKR, respectively.
2.4.2. Influenza virus PB2 627K modulates nucleocapsid detection by RIG-I

Adaptation of FLUAV to humans is required for establishment of an efficient virus infection. The influenza virus polymerase subunit PB2 has been described as a major determinant of host switching [74]. In particular, amino acid substitution at PB2 residue 627 from an avian E (glutamic acid) signature to mammalian adapted K (lysine) has been extensively characterized [120]. It was described that mammalian adaptation mutation PB2 627K results in an increased replication in mammalian cells whereas in chicken cells no major replication differences between avian PB2 627E and mammalian PB2 627K were detectable. Furthermore, different groups observed reduced interaction of avian PB2 627E with the viral nucleocapsid and this failure in nucleocapsid assembly was absent in chicken cells [66, 80, 96]. The molecular background for replication advantages of mammalian PB2 627K remained elusive [15, 74, 118]. Interestingly, chicken cells, where no major differences between avian PB2 627E and mammalian PB2 627K could be observed, lack RIG-I [4]. This raises the question whether adaptation mutation PB2 627K might be an influenza virus evasion strategy to avoid immediate RIG-I recognition in mammalian cells to enhance virus replication (manuscript 4.3, (Weber et al., in press)).

To address this question, the potential of the incoming nucleocapsids of four diverse influenza virus strains (A/quail/Shantou/2061/00 (H9N2), A/Thai/KAN-1/04 (H5N1), pandemic A/Hamburg/05/2009 (pH1N1), or A/WSN/33 (H1N1)) with either the avian PB2 627E or mammalian PB2 627K to promote RIG-I conformational switching was compared. Strikingly, nucleocapsids with the avian PB2 627E induced a robust RIG-I activation, which was strongly reduced in the presence of the mammalian signature PB2 627K (manuscript 4.3, Fig. 3A). Also by performing co-sedimentation assay, avian PB2 627E induced a more robust shift of RIG-I, TRIM25 and MAVS containing fractions in the gradient indicating an enhanced activation of the RIG-I signaling pathway in comparison to nucleocapsids bearing mammalian PB2 627K (manuscript 4.3, Fig. 3C).

Despite the clear effect on RIG-I activation, no significant differences between induction of the IFN response by an aborted replication cycle of avian PB2 627E or mammalian PB2 627K containing FLUAV were observed (manuscript 4.3, Fig. 3D). Likewise, the activation of the IFN transcription factor IRF3 by FLUAV nucleocapsids bearing either PB2 627E or PB2 627K did not differ. This indicates that mammalian adaptation mutation PB2 627K prevents RIG-I activation without major effects on IFN induction.

To test the influence of RIG-I on virus replication, wt (wildtype) or ΔRIG-I (RIG-I deficient) HEK293 cells were infected with A/Thai/KAN-1/04 (H5N1) with avian PB2 627E or mammalian PB2 627K signature. Allowing multi-cycle growth for 24 hours, titer differences between viruses with avian and mammalian polymerase complex were reduced from 50,000 fold in wt cells to 400 fold in ΔRIG-I cells (manuscript 4.3, Fig. 4D). These results could be further supported by monitoring the expression of viral nucleoprotein NP. Compared to wt cells, in ΔRIG-I cells viruses with the avian PB2 627E show an earlier NP synthesis, whereas viruses with the mammalian PB2 627K are not
Results

strongly affected by the presence or absence of RIG-I (manuscript 4.3, Fig. 4E, S4C and S4D). Likewise, in chicken DF-1 cells, naturally lacking RIG-I [4], FLUAV bearing the avian PB2 627E or mammalian PB2 627K express NP to a similar extent (manuscript 4.3, Fig. 4C, upper panel). However, overexpression of human RIG-I severely reduces NP expression of the avian PB2 627E virus, whereas replication efficiency of FLUAV with the mammalian PB2 627K is not strongly influenced (manuscript 4.3, Fig. 4C, middle panel).

As mentioned above, no striking differences in IFN induction stimulated by nucleocapsids bearing avian or mammalian polymerase complex could be observed (manuscript 4.3, Fig. 3D). Therefore, we wondered whether the IFN response has an influence on the different replication efficiencies of FLUAV with either avian PB2 627E or mammalian PB2 627K. Curiously, by using ΔMAVS (MAVS depleted) HEK293 cells incapable to transmit RIG-I signaling for IFN induction, no rescue of NP expression of FLUAV with avian PB2 627E could be detected (manuscript 4.3, Fig. 5A). This indicates that inhibition of RIG-I signaling seems to be not sufficient to regain replication potential of avian viruses. To support this hypothesis, ΔRIG-I cells or chicken DF-1 cells were transcomplemented with either RIG-I wt or signaling incompetent RIG-I K270A mutant. Strikingly, RIG-I K270A did impair PB2 627E virus as much as RIG-I wt (manuscript 4.3, Fig. 5B and Fig. 5C). Antiviral activity of RIG-I against nucleocapsids bearing the avian PB2 627E is consequently independent of RIG-I downstream signaling function. Thus, RIG-I binding to the avian nucleocapsids seems to be sufficient to impair virus replication suggesting a novel direct antiviral RIG-I activity.

Mammalian PB2 627K adaptation was described to enhance association of the polymerase complex with the nucleocapsid [66]. To test whether RIG-I affects association of the avian polymerase complex with the nucleocapsids in mammalian cells, co-IPs were performed. In wt cells the avian polymerase complex was less efficiently coprecipitated with the nucleocapsids whereas the PB2 627E-nucleocapsid interaction was slightly increased in ΔRIG-I cells (manuscript 4.3, Fig. 6A). By performing a RIG-I specific co-IP, more nucleocapsids with the avian PB2 627E could be coprecipitated in comparison to nucleocapsids bearing PB2 627K (manuscript 4.3, Fig. 6B). Although less nucleocapsids of a mammalian adopted strain could be precipitated, more PB2 627K was associated. This indicates that stronger binding of the mammalian polymerase complex to the 5’ppp panhandle prevents RIG-I recognition and hence displacement of the polymerase complex. To test this, a polymerase complex disassembling compound (PB1-T6Y) was applied [141]. Destabilization of the viral polymerase complex converts nucleocapsids with the mammalian PB2 627K to strong RIG-I activators (manuscript 4.3, Fig. 6D). However, the potential of PB2 627E bearing nucleocapsids to promote RIG-I conformational switching is more modestly affected. This implies that the strength of the polymerase complex-nucleocapsid interaction influences RIG-I activation.

In summary, we found that RIG-I acts as an influenza virus restriction factor by promoting the dissociation of the weakly interacting avian polymerase complex, thus impairing virus replication. The
inhibitory effect of RIG-I represents moreover a so far undescribed direct, signaling independent antiviral activity. On the other hand, influenza viruses have adapted to allow efficient replication in mammalian cells. Mammalian adaptation mutation PB2 627K enhances binding to the nucleocapsid, thereby preventing RIG-I recognition.

2.4.3. Lassa virus nucleoprotein promotes proteasomal degradation of PKR

Also arenaviruses need to adapt to evade immune recognition for a productive infection. It is known that arenaviruses have evolved diverse strategies to simultaneously attack diverse steps of RIG-I signaling [134, 135]. However, it remained elusive how arenaviruses prevent immediate detection by PKR. Interestingly, we observed that arenavirus nucleocapsids stimulated PKR phosphorylation and conformational switching only if the translation inhibitor CHX was applied (manuscript 4.4, Fig. 3B). This raises the question whether arenaviruses encode for a yet unidentified PKR antagonist. To investigate this hypothesis, we focused on LASV. By performing co-IP, an interaction of PKR with incoming (manuscript 4.4, Fig. 4A, right panel) and newly synthesized LASV nucleocapsids could be observed (manuscript 4.4, Fig. 4A, left panel). However, phosphorylated PKR could only be detected if protein synthesis was inhibited. As a control, a RIG-I specific co-IP was included. RIG-I does not interact with incoming LASV nucleocapsids (manuscript 4.4, Fig. 4A, left panel), which is in agreement with the results of Marq et al. that 5’overhang structures impair RIG-I recognition [75, 76]. A weak interaction of RIG-I with LASV nucleocapsids could only be observed if full replication cycle was allowed, which might reflect erroneous replication intermediates (manuscript 4.4, Fig. 4A, right panel). To identify the protein responsible for PKR inhibition, LASV NP (nucleoprotein), GP (glycoprotein) and Z (matrixprotein) were transfected with increasing concentrations together with PKR at a constant plasmid amount. Thereby, overexpression of PKR alone resulted in PKR activation due to proximity of the single PKR molecules allowing auto-phosphorylation. LASV GP and Z further increased the phosphorylation status of PKR whereas increasing amounts of LASV NP dramatically reduced PKR activation (manuscript 4.4, Fig. 4B). By performing co-IP, an interaction of LASV NP, but not GP or Z, with PKR was identified (manuscript 4.4, Fig. 4C). Thus, LASV NP is able to interact with PKR and impair PKR phosphorylation. To validate the involvement of the proteasome, the proteasomal inhibitor MG132 was tested. In absence of MG132, LASV NP overexpression results in decreased PKR phosphorylation as expected (manuscript 4.4, Fig. 4D, left panel). However, PKR activation in LASV NP-overexpressing cells can be rescued if the proteasomal degradation pathway is blocked (manuscript 4.4, Fig. 4D, right panel). Also during LASV infection MG132 application can prevent NP promoted PKR inhibition (manuscript 4.4, Fig. 4E). Surprisingly, MG132 treatment likewise rescued PKR activation in TCRV and JUNV infected cells.

In summary, we discovered that LASV NP interacts with PKR and promotes the proteasomal degradation of activated PKR. This seems to be a conserved mechanism within the arenavirus family.
since MG132 treatment rescues PKR activation during Old World LASV infection, but also in New World TCRV and JUNV infected cells.
3. Discussion

3.1. RIG-I as an immune sensor of incoming viral nucleocapsids

A rapid immune response to virus infection is crucial to ensure survival of the host. *In vitro* studies helped to identify major determinants triggering a RIG-I dependent antiviral IFN response, like dsRNA and a 5’ppp [46, 91, 111, 113]. However, in the natural context of infection the first viral structure able to stimulate RIG-I activation still needed to be resolved. Here, RIG-I is presented as an immune sensor of incoming bunyavirus and influenza virus nucleocapsids enabling an immediate antiviral response. RIG-I directly interacts with the incoming nucleocapsids independent of viral or cellular RNA synthesis or mammalian cofactors. RIG-I contacts the nucleocapsids via a single site, most likely via the 5’ppp dsRNA panhandle. Thereby, interaction is dependent on the presence of a 5’ppp and a dsRNA, since viral nucleocapsids lacking these features failed to mediate RIG-I activation. This is in contrast to the observation that 3’ untranslated regions of the influenza virus genome induce RIG-I activation independent of a 5’ppp [21]. However, these data are based on *in vitro* assays transfecting unpackaged viral or synthetic RNA into cells. During infection, RIG-I activation seems to be dependent on a 5’ppp. This was also proposed by the A.M. Pyle group, who likewise detected RIG-I activation in absence of a 5’ppp in *in vitro* experiments, but postulated that during virus infection the 5’ppp would be required for robust RIG-I binding and activation in presence of a less abundant agonist [60]. Indeed, our results demonstrate that during virus infection, an interaction of RIG-I with 5’ppp nucleocapsids stimulates conformational switching and oligomerization, two hallmarks of RIG-I activation. Activated RIG-I promotes then a signaling cascade that stimulates the activation of the IFN transcription factor IRF3 and eventually culminates in the induction of the antiviral IFN response. This defines the release of viral nucleocapsids as the first time-point of a RIG-I-dependent immune recognition in the natural context of virus infection. In former studies, transfected measles and vesicular stomatitis virus nucleocapsids could already be identified as immune stimulators [124, 125]. However, during transfection the correct assembly of the viral nucleocapsids cannot be ensured and furthermore, the immune response was not associated with RIG-I. Immune recognition of incoming nucleocapsids is, however, in contrast to previous reports suggesting that replication intermediates occurring later during viral infection cycle would trigger RIG-I-dependent immune responses [5, 101]. Especially for FLUAV, in a recent paper it was argued that the IFN induction is dependent on RNA synthesis [59, 86]. Killip *et al.* observed an activation of the IFN transcription factor IRF3 only if RNA synthesis and nuclear export of the progeny nucleocapsids was permitted [59]. However, we detected IRF3 activation even if the transcription inhibitor ActD was applied or nuclear export was blocked by LMB treatment. The discrepancy could be due to the chosen time-point of analysis. In case of Killip *et al.*, IRF3 translocation was validated eight hours post infection, a time-point where IRF3 might have already disappeared from the nucleus [59]. In our study, IRF3 activation was observed as early as one hour post-infection. This indicates
that also incoming FLUAV nucleocapsids serve as stimulators of the antiviral IFN response, as previously demonstrated for influenza B viruses [86].

It is somewhat intriguing how RIG-I gains access to the 5’ppp panhandle, which is covered by the viral polymerase. Recent structural data of the bat influenza virus panhandle further complicates the view of the accessibility of the 5’terminus [90]. The 5’terminus folds by intrastrand base-pairing into a hook-like structure followed by a dsRNA stretch of 14 base pairs. Moreover, the whole structure is deeply covered within the polymerase complex. It can be hypothesized that during the transit to the nucleus the association of the polymerase complex with the panhandle is dynamic and to some extend flexible. This is in agreement with the observation that the polymerase complex, especially PB2, is mobile within the structure [102]. A partial release of the 5’terminal hook may enable RIG-I binding to the 5’ppp. Because of the four base-paired nucleotides forming the hook structure, the nucleotide bearing the 5’ppp is base-paired enabling a robust interaction of the RIG-I CTD domain. Additionally, dsRNA stretches are associated with a stabilization of the RNA helix in a fixed conformation supporting engagement of the RIG-I helicase domain [62]. Due to the high affinity of RIG-I for the 5’ppp [128], RIG-I can rapidly entrap its agonist, even if it is only shortly exposed. RIG-I interaction with the 5’ppp may allow first structural rearrangements of RIG-I thereby triggering a disposition of the polymerase complex. This is supported by the hypothesis of Schmidt et al., who argue that RIG-I can drive an ATP-dependent removal of proteins [114]. It seems also plausible, that the panhandle is briefly exposed allowing RIG-I to rapidly interact. Both hypotheses, however, require further investigation.

In summary, RIG-I is in a yet to be identified mechanism able to engage the 5’ppp dsRNA panhandle of incoming nucleocapsids. This interaction induces RIG-I activation, enabling downstream signaling for induction of an immediate antiviral response.

3.2. PKR contributes to immediate pathogen recognition

Immune recognition of incoming nucleocapsids allows an immediate antiviral response. However, viruses have evolved to evade RIG-I recognition. Therefore, alternative immune receptors are required to ensure immediate induction of antiviral defense mechanisms or to support RIG-I immune recognition for a sustained response.

In the present thesis, we identified PKR involvement in immediate virus recognition. PKR interacts with the IGR of incoming bunyavirus RVFV S segmented nucleocapsids and incoming New World arenavirus TCRV and JUNV and Old World arenavirus LASV nucleocapsids. Like RIG-I, PKR interacts with the nucleocapsids only via a single contact site. By analyzing the predicted RVFV and arenavirus IGR, a hairpin structure can be observed with a central dsRNA stem with internal loops and sporadic mismatches. A 16 base pair dsRNA stem with additional 10 to 15 nucleotides of single-stranded tails was previously described as the minimal requirement for PKR activation [150]. Thereby, the dsRNA does not have to possess perfect base pairing since PKR can tolerate non-Watson-Crick
structures and also internal loops as long as the overall A-form geometry of the RNA is retained [7, 8]. Hence, the IGR of the RVFV small genome segment and all genome segments of TCRV, JUNV and LASV present all criteria of a potential PKR agonist. Previous studies demonstrated an interaction of PKR with structured elements of viral RNA, like Hepatitis C virus IRES (internal ribosomal entry site) [119] or dimerized stem-loop structures of TAR (transactivation RNA) of HIV [43].

In vitro experiments by Dauber et al. suggest that 5’ppp panhandle structures also serve as PKR agonists [19]. Our data indicates, however, that RIG-I, but not PKR, interacts with the 5’ppp dsRNA panhandle structure in the natural context of infection. Thereby, RVFV M segmented nucleocapsids presenting a 5’ppp dsRNA panhandle, but lacking the IGR promoted RIG-I conformational switching and failed to stimulate PKR activation. It can be postulated that RIG-I outcompetes PKR for 5’ppp panhandle interaction. Another plausible explanation could be that engagement of the 5’ppp terminus by PKR plays a role only during later time-points of infection, when misencapsidated side products of viral replication occur. So far, PKR was not associated with the ability to remove proteins from RNA as it was postulated for RIG-I [114]. Thus, PKR might not gain access to the panhandle of incoming nucleocapsids covered by the viral polymerase.

However, the question about the accessibility of the IGR within the nucleocapsid complex remains elusive. In case of orthobunyaviruses, like LACV, the RNA is deeply buried within the nucleocapsids [3] hindering PKR association. However, RVFV and arenavirus nucleocapsids are less tightly packaged leaving the viral genome partially accessible [40, 99, 100]. Exposure of the viral RNA alone is nevertheless not sufficient for PKR activation since RVF VLPs containing either M or S segment, with the same packaging grade of the viral RNA within the nucleocapsid, do only activate PKR if the IGR is present. Moreover, data by Moy et al. indicates that the IGR is exposed for host protein interaction [83]. The group thereby identified an interaction of human DDX17 or Drosophila Rm62 with the IGR of RVFV S segment indicating that also PKR could gain access to the IGR. Indeed, our data demonstrates that PKR is able to engage the IGR within the viral nucleocapsid in the natural context of virus infection. Protein interaction with the IGR was proposed to have a negative impact on virus replication [83]. It can therefore be hypothesized that PKR association with the RVFV and arenavirus IGR could also impair virus replication independent of its downstream signaling activity. This represents an exciting assumption, but still requires further investigation.

Taken together, PKR contributes to immediate virus recognition by interacting with the IGR of incoming viral nucleocapsids.

3.3. Immune evasion strategies to prevent immediate recognition

Antiviral immunity acts as a major barrier to virus infection, which must be circumvented to enable a productive infection. Viruses have therefore evolved to prevent immediate recognition and activation of antiviral defense mechanisms.
As presented above, incoming FLUAV nucleocapsids serve as immediate triggers of a RIG-I-dependent IFN response. To ensure an efficient replication cycle, influenza viruses have evolved multiple strategies to circumvent immune recognition by RIG-I, downstream signaling and activation of an antiviral response. On one hand, during evolution Orthomyxoviridae have located viral transcription and replication to the nucleus thereby limiting exposure of potential agonist to cytoplasmic immune sensors. Furthermore, immune recognition is actively impaired by sequestering dsRNA by viral NS1 [42] or dsRNA unwinding via the cellular helicases UAP56 and URH49 [136, 137]. Also FLUAV polymerase complex interferes with the host immune system. Thereby, MAVS downstream signaling ability is impaired by influenza virus polymerase subunits PB1, PB2 and PA [34, 53] and a variant of PB1-F2 [24, 127]. Like NS1 protein [49], the polymerase complex is already present within the virion allowing a rapid interference with immune recognition.

Here, we present a single substitution at position 627 of PB2 from avian adapted glutamic acid (E) signature to mammalian lysin (K) as a novel FLUAV strategy to impair immediate immune recognition by RIG-I. A change from avian PB2 627E to mammalian PB2 627K has been extensively characterized in the context of host adaptation [74]. Also adaptation to the host immune system is required to ensure efficient replication in a new host. Previous data suggested that adaptation mutation PB2 627K positively affects viral replication in mammalian cells, which was accompanied with increased nucleocapsid stability [66, 80, 96]. The PB2 627 position is in close proximity to the NP binding site and hence an unstable binding between PB2 627E and NP might lead to a failure in nucleocapsid assembly [80, 82].

Our data indicates that RIG-I specifically recognized and got activated by nucleocapsids with avian polymerase complex whereas nucleocapsids bearing mammalian PB2 627K failed to promote RIG-I activation. Furthermore, RIG-I specifically interfered with the association of the avian polymerase complex with the nucleocapsids. This is in agreement with previous reports describing a disturbed interaction of PB2 627E with the viral nucleocapsids in mammalian cells, but the responsible host factor or mechanisms was not identified [66, 80, 96]. In contrast, Cauldwell et al. argued that mammalian adaptation mutation PB2 627K does not enhance stability of the polymerase complex, but rather increases the amount of replicated RNA [14]. This would consequently lead to the false interpretation of co-IP data that more mammalian polymerase complex associated to the nucleocapsids excluding the fact that less avian nucleocapsids are present. However, in our case cells were treated with CHX and LMB to block viral replication and still, in presence of RIG-I, more mammalian PB2 627K was associated to the incoming nucleocapsids in comparison to avian PB2 627E. In RIG-I depleted cells, interaction of the avian PB2 627E bearing polymerase complex with the nucleocapsids was partially rescued. This implicates that RIG-I specifically affects the association of a destabilized avian polymerase complex. PB2 627K, however, enhances interaction of the polymerase complex with the nucleocapsid thus limiting RIG-I recognition of the incoming nucleocapsids. This is also supported
by the observation that disturbance of the polymerase complex assembly through applying competing peptides reverts PB2 627K nucleocapsids to a RIG-I stimulating agonist.

It was furthermore described that the mutation at position 627 of influenza virus PB2 does not alter the structure of the polymerase subunit, but rather disrupts a basic patch on the surface of the protein [122]. The change of surface charge might alter the association of the polymerase complex with host factors. Interaction with either enhancing or suppressing cellular proteins might therefore be responsible for the diverse polymerase activity between PB2 627E and 627K [96]. Indeed, the García-Sastre group described several proteins differentially regulating polymerase activity depending if there is an E or K at position 627 of PB2. Interestingly, DDX17, a DEAD box RNA helicase like RIG-I, was among the identified proteins [10]. In our hands, an interaction of RIG-I specifically with avian PB2 627E or the mammalian PB2 627K could not be detected. However, others did identify an association of FLUAV polymerase complex subunits with RIG-I [69]. Due to sensitivity problems, our experiments might have failed to detect this interaction. The biological significance of polymerase-RIG-I interaction still needs to be resolved. It was, however, demonstrated that the interaction does not affect RIG-I mediated IFN signaling [69]. It can be postulated that RIG-I entraps the viral polymerase complex to the panhandle promoter region and blocks polymerase processivity by interaction, since the RIG-I CTD is anchored via association to the 5’ppp. This might promote an IFN signaling-independent block of viral replication. Our data likewise indicate an IFN-independent mechanism of inhibiting virus replication of FLUAV with the avian signature. Firstly, by validating the ability of incoming nucleocapsids bearing either avian PB2 627E or mammalian PB2 627K to stimulate IFN induction no striking differences were observed. Furthermore, depletion of cells of the signaling molecule MAVS did not rescue replication of the FLUAV with avian PB2 627E. Likewise, a signaling-incompetent mutant RIG-I with an active binding capacity inhibited replication of FLUAV with avian PB2 627E as much as RIG-I wt did. Taken together, not the downstream IFN signaling ability of RIG-I promotes the antiviral effect against FLUAV with the avianized polymerase complex, but rather already the association with the panhandle promoter. This indicates a novel direct antiviral activity of RIG-I independent of its signaling ability.

In conclusion, RIG-I detects incoming influenza virus nucleocapsids. Activation of RIG-I is thereby enhanced in presence of an avian PB2 627E. Poor association of the polymerase complex with the nucleocapsid might allow full engagement of RIG-I with the 5’ppp panhandle promoting its activation. RIG-I interaction with the panhandle alone dislocates the avian polymerase and impairs virus replication independent of RIG-I mediated IFN signaling. However, a polymerase complex with the mammalian signature PB2 627K shows enhanced interaction with the nucleocapsid impairing RIG-I recognition. This suggests that also for recognizing other viral nucleocapsids the polymerase needs to be disposed for enabling RIG-I binding and activation. It would be hence interesting to investigate whether other viruses possess similar adaptation mutations to avoid immediate RIG-I recognition.
Despite RIG-I, also PKR contributes to immediate recognition of virus infection. PKR, however, detects the IGR of incoming nucleocapsids presented by viruses using ambisense coding strategy, like RVFV S and arenavirus genome segments. RVFV NSs promotes the proteasomal degradation of PKR thereby limiting PKR recognition and its antiviral activity [37]. An evasion strategy by arenavirus able to inhibit PKR activation was not identified so far.

Here, we present that arenaviruses promote, like RVFV, the proteasomal degradation of PKR. Specifically, LASV NP could be identified as the responsible PKR antagonist. LASV NP interacts with PKR and promotes PKR degradation via the proteasomal pathway in a dose dependent manner. However, NP encapsidating the viral genome did not affect the activation status of PKR indicating that only free LASV NP was comprising PKR antagonistic properties. LASV NP possesses the ability to bind dsRNA [39], which could serve as a scaffold for PKR interaction. Co-IP data of LASV proteins expressed in absence of viral RNA indicates, however, that the interaction of LASV NP with PKR is rather independent of viral RNA. Nevertheless, a complex formation of LASV NP with cellular RNA cannot be excluded. It was reported in this regard that recombinant nucleoproteins of NSV can associate with cellular RNA into nucleocapsid like structures [106]. Moreover, LASV NP comprises the ability to oligomerize [68]. Whether NP monomers or LASV oligomeric complexes promote PKR antagonism still needs to be clarified. It was, however, previously described that the NP oligomerization is no strict requirement for impairing the IFN response [68]. Hastie et al. reported that the dsRNA specific 3’ to 5’ exonuclease activity of LASV NP is essential for inhibiting IFN signaling [39]. Since incoming nucleocapsids do not present long dsRNA stretches to the host immune system, degradation of dsRNA might play a distinct immune evasion strategy during later infectious stages.

Arenavirus NP is well conserved among Old and New World arenaviruses, suggesting that an IFN antagonistic activity may be a shared feature [39, 40]. Indeed, lymphocytic choriomeningitis virus (Old World arenavirus) and Whitewater Arroyo virus, Pichinde virus, JUNV, Machupo virus and Latino virus, but not TCRV, (New World arenaviruses) were able to impair the IFN and proinflammatory response [77, 78, 104]. Likewise, arenaviruses use proteasomal degradation as a common PKR antagonism, but the responsible TCRV and JUNV protein inducing PKR inhibition remains elusive. TCRV, incapable to impair IFN signaling [78], inhibited PKR activation. This highlights the requirement to prevent early recognition by PKR to ensure efficient virus propagation, even in the case of apathogenic viruses like TCRV.

Despite of proteasomal degradation of PKR by arenaviruses, other viruses have also evolved strategies to prevent antiviral activity of PKR. Thereby, Kaposi-sarcoma herpesvirus vIRF2 and vaccinia virus K3L block PKR auto-phosphorylation [13, 105]. Hepatitis C virus E2 protein presents homologies with PKR and eIF2α and can serve as a pseudosubstrate [123]. This broad spectrum of viruses ranging from DNA to RNA viruses antagonizing PKR indicates the relevance of PKR inhibition for efficient virus replication.
In summary, we found that LASV NP and a yet to be identified TCRV and JUNV protein promote the proteasomal degradation of PKR. Thereby, LASV NP interacts with PKR most likely independent of viral RNA and induces PKR degradation via the proteasomal pathway. Therefore, arenavirus have evolved an efficient strategy to counteract immediate recognition and activation of antiviral defense mechanisms.

3.4. Concluding remarks

The presented data provides evidence that incoming bunyavirus, FLUAV and arenavirus nucleocapsids are rapidly recognized by RIG-I and PKR. RIG-I recognition is dependent on the presence of a 5’ppp dsRNA panhandle, whereas PKR detects the IGR of genome segments using ambisense coding strategy. Hence, the first host response to virus infection already occurs immediately after virus entry. Furthermore, we describe a novel antiviral activity of RIG-I that is independent of its signaling ability. In this regard, already binding of RIG-I to the 5’ppp panhandle limits FLUAV replication.

Viruses evade immune recognition by various mechanisms. Here, we present two diverse strategies of FLUAV and arenaviruses to prevent immediate detection by the host immune system. For FLUAV, the mammalian adaptation PB2 627K stabilizes the polymerase complex association with the nucleocapsids, thus preventing RIG-I recognition of the 5’ppp dsRNA panhandle. In case of arenaviruses, we show that LASV NP prevents PKR sensing of the incoming nucleocapsids via induction of the proteasomal degradation of PKR. This later seems to be common within the arenavirus family since not only Old World arenavirus LASV, but also New World TCRV and JUNV induce PKR degradation via the proteasomal pathway.

It is known that the constant immune pressure leads to positive selection of viruses combining potent immune evasion with efficient replication strategies within the host cell. FLUAV bearing PB2 627K is associated with enhanced pathogenicity [120], like LASV being the causative agent of fatal human hemorrhagic fever [79]. This supports the idea that viral proteins involved in circumventing RLR and IFN signaling represent also determinants of virulence and pathogenesis [12, 129].

In conclusion, we pin down entry of viral nucleocapsids as the first time-point of immune recognition in the natural context of virus infection and give further insights how viruses have evolved to counteract immediate detection.

Despite these achievements, future studies are required to comprehend the complexity of virus-host interactions. It remains to be clarified how RIG-I gains access to the 5’ppp dsRNA panhandle covered by the viral polymerase and nucleoproteins. Furthermore, the direct antiviral activity of RIG-I combined with the fact that cofactors are not essential for RIG-I activation raises the question whether RIG-I would also perform an antiviral activity in the nucleus. Indeed, RIG-I was previously observed in the nucleus [69], however, no antiviral activity was associated. Interestingly, it was hypothesized
that an unknown factor interacts with avian PB2 627E in the nucleus, thereby limiting mobility and disturbing nucleocapsid assembly [28]. Whether RIG-I represents this factor is a tempting assumption. For FLUAV, several adaptation mutations associated with increased pathogenesis were described [15]. The effect of these mutations on cellular defense mechanisms was not addressed so far. It would be thus of interest to validate the influence of other adaptive mutations, despite PB2 627K, on the host immune response.

To effectively control virus infections, the generation of new therapeutically tools is required. Interestingly, peculiar poly-C (viral sense) and poly-G (viral anti-sense) repeats within the IGR were previously described as essential transcription termination signals [1]. These structures are therefore less likely to be altered by the virus despite the existing immune pressure. Moreover, interaction of a protein with the IGR of viral genomes was associated with reduced viral replication [83]. Thus, targeting of this conserved region, by for instance aptamers, might be a promising therapeutic approach to limit virus replication and ensure survival of the host.

Conclusively, detailed knowledge how viruses interfere with the RLR detection and signaling system may provide not only new insights into RLR signaling, but also virus pathogenesis itself. A better understanding of the pathogenic potential of viruses would thereby allow to rapidly react and provide preparedness for future epidemics or even pandemics.

Immediate virus recognition by RIG-I and PKR and viral counterstrategies.

Exemplarily, immune recognition of incoming influenza A virus (FLUAV) and Lassa virus (LASV) nucleocapsids by RIG-I and PKR is shown, respectively. RIG-I rapidly detects the 5’ppp dsRNA panhandle upon release of FLUAV nucleocapsids
into the host cytoplasm. PKR, however, reacts to the intergenic region of incoming nucleocapsids. To antagonize immediate immune recognition, FLUAV mammalian adaptation mutation PB2 627K stabilizes association of the polymerase complex with the nucleocapsids, thereby preventing RIG-I recognition. For PKR evasion, LASV nucleoprotein (NP) interacts with PKR and promotes its degradation via the proteasome.
4. Original publications and manuscripts

4.1. Monitoring activation of the antiviral pattern recognition receptors RIG-I and PKR by limited protease digestion and native PAGE

Own contribution:

I established the techniques in our lab and performed all experiments shown in Fig. 1 and 2. I contributed to writing of the manuscript.

______________________________
Michaela Gerlach
Monitoring Activation of the Antiviral Pattern Recognition Receptors RIG-I And PKR By Limited Protease Digestion and Native PAGE

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Abstract

Host defenses to virus infection are dependent on a rapid detection by pattern recognition receptors (PRRs) of the innate immune system. In the cytoplasm, the PRRs RIG-I and PKR bind to specific viral RNA ligands. This first mediates conformational switching and oligomerization, and then enables activation of an antiviral interferon response. While methods to measure antiviral host gene expression are well established, methods to directly monitor the activation states of RIG-I and PKR are only partially and less well established.

Here, we describe two methods to monitor RIG-I and PKR stimulation upon infection with an established interferon inducer, the Rift Valley fever virus mutant clone 13 (Cl 13). Limited trypsin digestion allows to analyze alterations in protease sensitivity, indicating conformational changes of the PRRs. Trypsin digestion of lysates from mock infected cells results in a rapid degradation of RIG-I and PKR, whereas Cl 13 infection leads to the emergence of a protease-resistant RIG-I fragment. Also PKR shows a virus-induced partial resistance to trypsin digestion, which coincides with its hallmark phosphorylation at Thr 446. The formation of RIG-I and PKR oligomers was validated by native polyacrylamide gel electrophoresis (PAGE). Upon infection, there is a strong accumulation of RIG-I and PKR oligomeric complexes, whereas these proteins remained as monomers in mock infected samples.

Limited protease digestion and native PAGE, both coupled to western blot analysis, allow a sensitive and direct measurement of two diverse steps of RIG-I and PKR activation. These techniques are relatively easy and quick to perform and do not require expensive equipment.

Introduction

A crucial event in antiviral host defense is the rapid detection of the pathogen by the so-called pattern recognition receptors (PRRs)¹,². Intracellular detection of RNA virus infection is dependent on two cytoplasmic RNA helicases, RIG-I (retinoic acid inducible gene I) and MDA5 (melanoma differentiation associated protein 5)³-⁵. RIG-I is composed of two N-terminal caspase recruitment domains (CARDs), a central DECH-box type RNA helicase domain, and a C-terminal domain (CTD)⁴,⁶. Whereas the CTD and the helicase domain are required for recognition of non-self (viral) RNAs, the CARDs mediate downstream signaling leading to establishment of an antiviral host status.

If RIG-I is in the silent state, i.e. in the absence of a specific RNA ligand, the second CARD interacts with the central helicase domain and keeps RIG-I in an auto-inhibitory conformation⁷-¹¹. RIG-I binds to short double-strand (ds) RNA bearing a 5'-triphosphate (5'PPP), long dsRNA, and polyU/UUC-rich RNA, classic signature structures which are present on the genomes of many RNA viruses¹²-¹⁶. Two major characteristics of RIG-I activation are a switch to a closed conformation⁶,¹⁷ and the homo-oligomerization²,¹⁸,¹⁹. The conformational switch enhances RNA binding, exposes the CARDs for downstream signaling, and reconstitutes an active ATPase site²⁰. The formation of oligomeric RIG-I complexes leads to enhanced recruitment of downstream signaling adaptor molecules to form a platform for antiviral signal transduction²¹. The RIG-I-regulated signaling chain eventually activates the transcription factor IRF-3 for up-regulation of interferon (IFN-alpha/beta) genes and hence the gene expression of interferon stimulated genes (ISGs) for a full antiviral response²²-²⁴. One of the best characterized ISGs is the RNA-activated protein kinase (PKR)²⁵. PKR belongs to the family of eukaryotic translation initiation factor 2 alpha (eIF2α) kinases and is composed of an N-terminal double-stranded RNA binding domain and a C-terminal kinase domain. The kinase domain constitutes the dimerization interface crucial for PKR activation and carries out the catalytic functions of the protein. Binding of PKR to viral dsRNA leads to its conformational change permitting dimerization and auto-phosphorylation at Thr 446 among other residues. PKR then mediates phosphorylation of eIF2α, thereby blocking the translation of viral mRNAs²⁶-²⁷.

Both RIG-I and PKR undergo major structural rearrangements, form oligomeric complexes and are post-translationally modified by phosphorylation/dephosphorylation and ubiquitination²⁸-²⁹. For a better understanding of which viral RNA structures are activating RIG-I and PKR (and at what stage viral antagonists could be interfering), it is important to precisely determine the activation status. For both PRRs it was previously described that activation leads to the emergence of trypsin-resistant protein fragments²⁶,²⁷,³⁰ and higher-order
1. Seeding of A549 Cells for Infection

1. Cultivate a T75 flask of A549 cells at 37 °C and 5% CO₂ in cell culture medium (DMEM supplemented with 10% FCS, 526.6 mg/l L-glutamine, 50,000 U/l penicillin, and 50 mg/l streptomycin).

2. Before starting to harvest the cells, warm up cell culture medium, PBS and 0.05% trypsin-EDTA in a waterbath heated to 37 °C.

3. Remove the medium and wash the cells with 10 ml PBS. Remove the PBS again.

4. Add 3 ml of trypsin-EDTA and distribute equally in the flask. Transfer the flask in an incubator with 37 °C and 5% CO₂.

5. When all cells are detached, add 7 ml of cell culture medium, resuspend the cells, and transfer the cell suspension into a 15 ml Falcon tube.

6. Centrifuge the cells at 800 x g for 5 min at RT, remove the supernatant and resuspend the pellet in 10 ml fresh cell culture medium.

7. Count the cells with a counting chamber.

8. Add 2.5 x 10⁶ cells in 5 ml of cell culture medium in two T25 flasks each. Incubate for 16 hr at 37 °C and 5% CO₂. One flask serves for the mock control and one for Cl 13 infection.

2. Infection with Rift Valley Fever Virus Clone 13 (Cl 13)

1. NOTE: Cl 13 is an attenuated virus mutant which in Germany can be handled under BSL-2 conditions. Please refer to relevant national guidelines. Other typical IFN inducers would be Sendai virus (strain Cantell) or Newcastle disease virus.

2. Pre-warm PBS, serum-free medium, and cell culture medium containing 5% FCS.

3. Prepare 1.25 x 10⁷ PFU/ml of Cl 13 in serum-free medium to infect 2.5 x 10⁶ A549 cells with a multiplicity of infection (MOI) of 5. Prepare a slightly (roughly 10%) greater amount than needed to account for pipette errors.

4. Wash the cells with PBS as described under 1.3.

5. When all cells are detached, add 7 ml of cell culture medium, resuspend the cells, and transfer the cell suspension into a 15 ml Falcon tube.

6. Centrifuge the cells at 800 x g for 5 min at RT, remove the supernatant and resuspend the pellet in 10 ml fresh cell culture medium.

7. Count the cells with a counting chamber.

8. Add 2.5 x 10⁶ cells in 5 ml of cell culture medium in two T25 flasks each. Incubate for 16 hr at 37 °C and 5% CO₂. One flask serves for the mock control and one for Cl 13 infection.

3. Preparation of Cell Lysates

1. Prepare PBS / 0.5% Triton X-100 at 4 °C. Do not add serine protease inhibitors.

2. Wash the cells with cold PBS and add 10 ml of fresh PBS.

3. Scrape the cells off, transfer the cell suspension in a Falcon tube, and centrifuge at 800 x g for 5 min at RT.

4. Remove the supernatant and resuspend the cell pellet in 30 µl PBS / 0.5% Triton X-100. Transfer the lysate into a fresh 1.5 ml tube and incubate for at least 10 min at 4 °C.

5. Centrifuge the lysate at 10,000 x g for 10 min at 4 °C and transfer the clarified cell lysate (supernatant) into a fresh tube.

6. Determine the protein concentration by Bradford assay as described elsewhere²⁹.

7. Store at -20 °C or proceed to trypsin digestion (4.1) or native PAGE (5.1).

4. Determination of Conformational Changes of Pattern Recognition Receptors

1. TPCK-trypsin Treatment of Cell Lysates

   1. Dilute L-1-tosylamido-2-phenylethyl chloromethyl ketone-treated (TPCK) trypsin in PBS to a final working concentration of 2 µg/µl.

   2. Adjust in two new tubes a final protein concentration of 25 µg of each protein lysate (mock or Cl 13) in a final volume of 9 µl with PBS. Hence, it should be four tubes with 25 µg lysate each, two times mock and two times Cl 13 infection. One set as input control (untreated) and one set with treatment with TPCK-trypsin.

   3. Add 1 µl of PBS (untreated) or 1 µl of 2 µg/µl TPCK-trypsin (final concentration: 0.2 µg/µl) to the cell lysates and mix the reactions by pipetting. DO NOT freeze and thaw TPCK-trypsin aliquots, because it will compromise the efficiency of digestion.

   4. Incubate the lysates at 37 °C for 25 min. Stop the reaction by adding 5x denaturing sample buffer (250 mM Tris-HCl pH 6.8, 10% SDS, 50% glycerol, 25% β-mercaptoethanol, 0.5% bromphenol blue) and by boiling for 5 min at 95 °C. It is important to NOT extend the trypsin incubation time. In case no protease-resistant fragments are detectable, the time of trypsin digestion must be shortened.

   5. After boiling, the samples can be stored at -20 °C.

2. SDS Polyacrylamide Gel Electrophoresis (PAGE) and Western Blotting

   1. Load the samples on a sodium dodecyl sulfate (SDS) polyacrylamide gel containing a 5% stacking over a 12% resolving gel. Separate the proteins at 25 mA per gel until the bromphenol blue runs out.

   2. Activate a polyvinylidene fluoride (PVDF) membrane for 30 sec with methanol and put it into transfer buffer (48 mM Tris, 39 mM glycine, 1.3 mM SDS, 20% methanol).

   3. Transfer the proteins by semi-dry electroblotting, using a VELVET apparatus (Schleicher & Schuell) as described in ref 31,32.

   4. After the transfer, the PVDF membrane is blocked with 5% non-fat milk in PBS-T for 1 hr. To be used only the last Pipes/Triton X-100 wash.

   5. After blocking, the PVDF membrane is incubated with monoclonal antibodies against RIG-I and PKR for 1 hr at 37 °C.

   6. The PVDF membrane is washed with PBS-T and incubated with horseradish peroxidase (HRP) conjugate for 1 hr at 37 °C.

   7. The PVDF membrane is washed with PBS-T and immersed in a solution containing 50 mM Tris-HCl pH 7.5, 100 mM NaCl, 0.1% Tween 20, 0.05% SDS, and 10% bovine serum albumin.

   8. After incubation, the PVDF membrane is washed with PBS-T and immersed in a solution containing 50 mM Tris-HCl pH 7.5, 100 mM NaCl, 0.1% Tween 20, 0.05% SDS, and 10% bovine serum albumin.

   9. The PVDF membrane is washed with PBS-T and immersed in a solution containing 50 mM Tris-HCl pH 7.5, 100 mM NaCl, 0.1% Tween 20, 0.05% SDS, and 10% bovine serum albumin.

   10. After incubation, the PVDF membrane is washed with PBS-T and immersed in a solution containing 50 mM Tris-HCl pH 7.5, 100 mM NaCl, 0.1% Tween 20, 0.05% SDS, and 10% bovine serum albumin.

   11. After incubation, the PVDF membrane is washed with PBS-T and immersed in a solution containing 50 mM Tris-HCl pH 7.5, 100 mM NaCl, 0.1% Tween 20, 0.05% SDS, and 10% bovine serum albumin.

   12. After incubation, the PVDF membrane is washed with PBS-T and immersed in a solution containing 50 mM Tris-HCl pH 7.5, 100 mM NaCl, 0.1% Tween 20, 0.05% SDS, and 10% bovine serum albumin.

   13. After incubation, the PVDF membrane is washed with PBS-T and immersed in a solution containing 50 mM Tris-HCl pH 7.5, 100 mM NaCl, 0.1% Tween 20, 0.05% SDS, and 10% bovine serum albumin.

   14. After incubation, the PVDF membrane is washed with PBS-T and immersed in a solution containing 50 mM Tris-HCl pH 7.5, 100 mM NaCl, 0.1% Tween 20, 0.05% SDS, and 10% bovine serum albumin.

   15. After incubation, the PVDF membrane is washed with PBS-T and immersed in a solution containing 50 mM Tris-HCl pH 7.5, 100 mM NaCl, 0.1% Tween 20, 0.05% SDS, and 10% bovine serum albumin.

   16. After incubation, the PVDF membrane is washed with PBS-T and immersed in a solution containing 50 mM Tris-HCl pH 7.5, 100 mM NaCl, 0.1% Tween 20, 0.05% SDS, and 10% bovine serum albumin.

1. After boiling, the samples can be stored at -20 °C.

1. Dilute L-1-tosylamido-2-phenylethyl chloromethyl ketone-treated (TPCK) trypsin in PBS to a final working concentration of 2 µg/µl.

2. Adjust in two new tubes a final protein concentration of 25 µg of each protein lysate (mock or Cl 13) in a final volume of 9 µl with PBS. Hence, it should be four tubes with 25 µg lysate each, two times mock and two times Cl 13 infection. One set as input control (untreated) and one set with treatment with TPCK-trypsin.

3. Add 1 µl of PBS (untreated) or 1 µl of 2 µg/µl TPCK-trypsin (final concentration: 0.2 µg/µl) to the cell lysates and mix the reactions by pipetting. DO NOT freeze and thaw TPCK-trypsin aliquots, because it will compromise the efficiency of digestion.

4. Incubate the lysates at 37 °C for 25 min. Stop the reaction by adding 5x denaturing sample buffer (250 mM Tris-HCl pH 6.8, 10% SDS, 50% glycerol, 25% β-mercaptoethanol, 0.5% bromphenol blue) and by boiling for 5 min at 95 °C. It is important to NOT extend the trypsin incubation time. In case no protease-resistant fragments are detectable, the time of trypsin digestion must be shortened.

5. After boiling, the samples can be stored at -20 °C.
3. Prepare the blotting with a semidy blotting system and allow the transfer of the proteins at 10 V for 1 hr. Take the membrane out, rinse it briefly with water, and let it dry.

4. Reactivate the membrane by shortly transferring into methanol. Wash for 5 min with TBS. Block with 10% skim milk in TBS for 1 hr at RT or at 4 °C O/N. Wash the membrane 3x for 5 min each with TBS.

5. Prepare antibody dilution as recommended in table 1 and incubate the membrane for 1 hr at RT or at 4 °C O/N.

6. Wash the membrane 3x for 10 min each with TBS-T. Add the appropriate secondary antibody coupled with horseradish peroxidase at a 1:20,000 dilution in 1% skim milk in TBS. Incubate 45 min at RT.

7. Wash the membrane 3x for 10 min each with TBS-T, and one additional time with TBS.

8. For signal detection use a commercial chemiluminescence kit and a digital gel imaging system.

3. Coomassie Brilliant Blue G-250 Staining

1. Perform SDS-PAGE as described in 4.2.1. Load samples on an SDS polyacrylamide gel and run the gel at 25 mA per gel until bromphenol blue runs out.

2. Perform Coomassie Brilliant Blue G-250 staining at RT. Do all incubations under constant shaking.

3. Transfer the gel to the fixation solution containing 40% methanol and 10% acetic acid for 30 min.

4. Exchange the buffer to destaining solution (25% ethanol and 8% acetic acid) and incubate for 5 min.

5. Stain the gel with 0.2% Coomassie Brilliant Blue G-250 in 40% methanol and 10% acetic acid for 1 hr.

6. Destain the gel with the destaining solution and exchange the buffer after 10 min, 30 min, and 60 min.

7. Store the gel in 25% ethanol, 8% acetic acid, and 4% glycerol at 4 °C.

8. Perform imaging and analysis as described under 4.2.8.

5. Analysis of Oligomeric States of Pattern Recognition Receptors

1. Native PAGE

   1. Prepare 50 µg of cell lysate in a final volume of 10 µl with PBS and add 5× native sample buffer (250 mM Tris-HCl pH 6.8, 1% sodium deoxycholate, 50% glycerol, 0.5% bromphenol blue) to a final concentration of 1x.

   2. Load the samples IMMEDIATELY on a native polyacrylamide gel with 5% as stacking and 8% as resolving gel. Any delay will result in a loss of native complexes.

   3. Run the gel at 20 mA per gel at 4 °C with 50 mM Tris-NaOH pH 9.0, 384 mM glycin as anode and 50 mM Tris pH 8.3, 384 mM glycin, 1% sodium deoxycholate as cathode buffer. After 1.5-2 hr (bromphenol blue band has left the gel approximately 45 min earlier) the electrophoresis is finished.

2. Western Blotting

   1. Activate a polyvinylidene fluoride (PVDF) membrane for 30 sec with methanol and put it into Towbin buffer (25 mM Tris, 192 mM glycin, 0.1% SDS, 20% methanol).

   2. Assemble a wet blot chamber according to the manufacturer’s instructions and fill the tank with Towin buffer.

   3. Perform the blotting with 250 mA for 1.5 hr at 4 °C.

   4. When blotting is finished proceed as described from 4.2.3 on.

### Representative Results

Recognition of a viral agonist by RIG-I or PKR triggers conformational switching and oligomerization. We assayed these two activation markers by limited protease digestion and native polyacrylamide gel electrophoresis (PAGE), respectively.

Human A549 cells were infected with Rift Valley fever virus clone 13 (Cl 13), which is characterized by a mutation of the IFN antagonist NSs. Due to the absence of functional NSs, Clone 13 strongly induces RIG-I and PKR, leading to the establishment of a robust antiviral state in the cells.

Trypsin digestion of mock infected cell lysates results in a rapid degradation of RIG-I, whereas Cl 13 infection leads to the generation of a 30 kDa resistant RIG-I fragment. Also PKR shows partial resistance to trypsin digestion in infected samples, which coincides with its phosphorylation. To monitor efficiency and specificity of trypsin digestion, gels were stained with Coomassie Brilliant Blue G-250. Untreated samples show equal amounts of loaded proteins. Subjecting mock and Cl 13 cell lysates to trypsin digestion leads to a comparable decrease of global protein amounts. This demonstrates that trypsin treatment has the same efficiency for mock and Cl 13-infected samples.

The formation of oligomeric complexes was assayed by native PAGE. In uninfected cells, only monomers of RIG-I and PKR were detected. As additional control we included the transcription factor IRF-3, which is present as a monomer but known to dimerize upon activation via e.g., RIG-I. Cl 13 infection leads to a strong accumulation of RIG-I oligomeric complexes in form of a smear and of PKR and IRF-3 dimers/oligomers as a defined protein band.

These results demonstrate that limited trypsin digestion and native PAGE are useful tools to monitor conformational changes and oligomer formation of RIG-I and PKR upon infection.
Figure 1. Conformational switch of RIG-I and PKR. A549 cells were mock infected or infected with Cl 13 at an MOI of 5. After 5 hr, cells were lysed in PBS supplemented with 0.5% Triton X-100 and cleared cell lysates were either left untreated or treated with trypsin. Samples were subjected to SDS PAGE followed by Western blotting (A) or Coomassie staining (B). Blots were stained against RIG-I, PKR, phosphorylated PKR (Thr 446) and against RVFV nucleoprotein (RVFV N) and beta-actin as infection and loading control, respectively. Please click here to view a larger version of this figure.
Discussion

Sensing the presence of viruses and activation of the antiviral type I IFN system are crucial for successful innate immune responses. Virus detection is thereby mediated by pathogen recognition receptors (PRRs) like RIG-I and PKR, enabling a rapid response and activation of antiviral defense mechanisms. Here, we describe two methods to directly evaluate the activation status of RIG-I and PKR.

Limited protease digestion as a tool to monitor conformational changes of RIG-I and PKR was first described by the groups of M. Gale Jr. and T. Fujita, and J. L. Cole, respectively. It represents a sensitive method to evaluate sensitivity alterations to trypsin treatment caused by conformational changes. Applying trypsin digestion, we detected rapid degradation of RIG-I and PKR in mock infected samples, whereas trypsin treatment of virus-infected cell lysates led to trypsin resistant RIG-I fragments. Similarly, a trypsin resistant PKR fragment was detected upon Cl 13 infection. This was accompanied by PKR phosphorylation at Thr 446, a widely used marker of PKR activation. Comparing trypsin digestion of...
mock and Cl 13 cell lysates by Coomassie staining demonstrates a comparable decrease of global protein levels. This indicates that formation of resistant fragments is specific for proteins like RIG-I and PKR.

The formation of oligomeric complexes of RIG-I, PKR and IRF-3 was monitored by native PAGE. Subjecting Cl 13-infected cell lysates to native PAGE, we detected RIG-I, PKR and IRF-3 oligomers, whereas these proteins remained as monomers in mock infected samples. RIG-I needs to form oligomers to activate downstream pathways. It was hypothesized that RIG-I oligomerization supports recruitment of cofactors to form a signaling platform for antiviral response mechanisms. The function of PKR dimerization is not entirely understood. Most likely, the PKR subunits in the dimer phosphorylate each other. The type I IFN system is tightly regulated on a transcriptional level, and IRF-3 represents one central transcription factor for the induction of IFN and ISGs. Central hallmarks of activation are phosphorylation, dimerization, and translocation to the nucleus, where it recruits the transcriptional co-activators p300 and CREB-binding protein (CBP) to initiate IFN mRNA synthesis. Analysis of IRF-3 dimerization is a widely used tool to monitor activation of the type I IFN response. Therefore, we used the detection of IRF-3 oligomeric complexes as a proof of principle for the oligomerization assays of RIG-I and PKR oligomerization. Indeed, allowing separation of protein lysates under non-denaturing conditions, we were able to detect IRF-3 dimerization in Cl 13 infected samples.

Undoubtedly, each lab will need to optimize the protocols of limited protease digestion and native PAGE. If confronted with no detection of any resistant proteins or too many resistant fragments, one might shorten or prolong the time of digestion, respectively. Differences can also be due to the specific TPCK-trypsin stock, as preparations slightly differ. Therefore, various TPCK-trypsin concentrations should be tested for optimization. Cell lysates can be prepared from other cell lines than A549, but adaptations of the protocol might be required. It is recommended to adjust the amount of total proteins for trypsin digestion and native PAGE according to the expression level of the protein of interest in this specific cell type. Moreover, limited detection or weak separation of oligomeric complexes by native PAGE can have several reasons and can be addressed as followed: make sure that the equipment is cleaned to remove remaining denaturing agents, keep all samples and the gel during the PAGE at 4 °C, and do not extend the time between sample preparation and loading onto the gel. Limited protease digestion and native PAGE have also been employed to monitor the activation of another important, closely related PRR, MDA5. Monitoring of MDA5 activation required different experimental conditions compared to RIG-I and PKR, and was therefore not included here.

In summary, point-by-point protocols for two useful and sensitive methods to measure activation of RIG-I and PKR are presented. Limited protease digestion and native PAGE, both coupled to Western blot analysis, permit monitoring of conformational changes and oligomerization, respectively. Using these methods, we had previously shown that RIG-I can be activated by nucleocapsids of various viruses directly after their entry into the cells.

The exact nature and origin of the RNA species relevant for RIG-I and PKR activation in infected cells are still not entirely solved. Furthermore, many viruses interfere with the functions of PRRs by a wide variety of strategies. The presented techniques enlarge the spectrum of methods by allowing an easy and direct measurement of RIG-I and PKR activation, are quick to perform, and do not require expensive equipment.

Disclosures

No conflicts of interest declared.

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References

4.2. Incoming RNA virus nucleocapsids containing a 5'-triphosphorylated genome activate RIG-I and antiviral signaling

Own contribution:

I performed experiments of Fig. 2B, Fig. 3, Fig. 4, Fig. 5 (whereas RIG-I constructs were cloned by Anna Mareike Schmidt), Fig. 6 and Fig. 7 and Suppl. Fig. 1, Suppl. Fig. 2H, Suppl. Fig. 3A-G, Suppl. Fig. 4, Suppl. Fig. 5 and Suppl. Fig. 6. I contributed to writing of the manuscript.

Michaela Gerlach
Incoming RNA Virus Nucleocapsids Containing a 5′-Triphosphorylated Genome Activate RIG-I and Antiviral Signaling

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SUMMARY

Host defense to RNA viruses depends on rapid intracellular recognition of viral RNA by two cytoplasmic RNA helicases: RIG-I and MDA5. RNA transfection experiments indicate that RIG-I responds to naked double-stranded RNAs (dsRNAs) with a triphosphorylated 5′ (5′ppp) terminus. However, the identity of the RIG-I stimulating viral structures in an authentic infection context remains unresolved. We show that incoming viral nucleocapsids containing a 5′ppp dsRNA “panhandle” structure trigger antiviral signaling that commences with RIG-I, is mediated through the adaptor protein MAVS, and terminates with transcription factor IRF-3. Independent of mammalian cofactors or viral polymerase activity, RIG-I bound to viral nucleocapsids, underwent a conformational switch, and homo-oligomerized. Enzymatic probing and superresolution microscopy suggest that RIG-I interacts with the panhandle structure of the viral nucleocapsids. These results define cytoplasmic entry of nucleocapsids as the proximal RIG-I-sensitive step during infection and establish viral nucleocapsids with a 5′ppp dsRNA panhandle as a RIG-I activator.

INTRODUCTION

Host defenses to RNA viruses are dependent on rapid detection by pathogen recognition receptors (PRRs). Intracellular recognition of virus infection is mediated by two cytoplasmic RNA helicases: RIG-I and MDA5 (termed RIG-like receptors, RLRs) (Kato et al., 2011). The binding of an RNA ligand to these PRRs leads to phosphorylation and dimerization of interferon regulated factor 3 (IRF-3), which subsequently activates genes for type I interferons (IFN-α/β) (Hiscott, 2007). These cytokines trigger the expression of IFN-stimulated gene (ISG) products that have antiviral and immunomodulatory activities (Randall and Goodbourn, 2008).

The infection cycle of RNA viruses consists of the phases attachment, entry, messenger RNA (mRNA) transcription, genome replication, assembly, and exit. Negative-strand RNA viruses carry an RNA-dependent RNA polymerase (RdRp) within their particles and immediately start transcribing their genome after entering the cell. The protein products of this so-called primary transcription then drive the replication of the genome via a positive-sense intermediate. Positive-strand RNA viruses do not carry an RdRp in their particles and directly translate their genome after entry. The newly synthesized RdRp then produces a negative-sense intermediate, mRNA, and progeny genome. Usually, the genome of both negative- and positive-strand RNA viruses is packaged by a viral nucleocapsid protein (often called N).

Most studies on activation of RLRs were based either on infection with RNA viruses undergoing a full replication cycle or on transfection of cells with naked viral or synthetic RNAs. The infection experiments established that RIG-I and MDA5 recognize mostly nonoverlapping subsets of viruses (McCartney and Colonna, 2009). The RNA transfection experiments revealed that RIG-I responds to long double-stranded RNA (dsRNA) molecules, short dsRNAs with a triphosphorylated 5′ (5′ppp) terminus, and poly-U/UC-rich sequences, whereas MDA5 activation is more dependent on branched dsRNA structures (Binder et al., 2011; Hornung et al., 2006; Kato et al., 2008; Pichlmair et al., 2006, 2009; Saito et al., 2008; Schlee et al., 2009; Schmidt et al., 2009). Despite these achievements, the question...
of the natural RLR ligands (i.e., which viral structures are stimulating the RLRs in the authentic infection context) is largely unsolved. Two recent RNA fractionation studies showed that full-length and shortened RNAs arising during genome replication of 5′ppp-RNA viruses are natural stimulators of RIG-I (Baum et al., 2010; Rehwinkel et al., 2010). However, it remained open whether these RIG-I ligands were naked RNA side products of viral genome replication or whether RIG-I could also recognize nucleoprotein-encapsidated RNA, the main viral structure in the infected cell. Our study presented here addresses this problem and indicates that RIG-I is capable of reacting to incoming, encapsidated RNA virus genomes. This immediate early IFN response requires the viral 5′ppp dsRNA “panhandle” structure but is independent of viral RNA synthesis. RIG-I thereby directly interacts with the panhandle on the viral nucleocapsids, switches conformation, oligomerizes, and triggers the activation of IRF-3.

RESULTS

Our first aim was to identify the earliest infection step that triggers IFN induction. In the hope of drawing conclusions on the responsible viral determinant, we employed a set of RNA viruses with different genomic features. Dependent on the particular virus, a block of viral mRNA translation by cycloheximide (CHX) has different effects on primary transcription and genome replication (Table S1 available online).

Entry of Negative-Sense RNA Viruses Can Activate IFN Induction in the Absence of Replication

Influenza A virus (FLUAV) and vesicular stomatitis virus (VSV) are negative-strand RNA viruses. Application of CHX allows particle attachment, entry of nucleocapsids, and primary transcription, but not genome replication (Figure S1). We measured the virus-inducible genes for IFN-β and ISG56 in human AS49 cells by real-time RT-PCR. Figure 1A shows that both FLUAV and VSV are strongly activating these genes, even when CHX was applied. Thus, an IFN response can occur before the viruses start replicating their genome.

To narrow down the IFN-relevant infection step, we repeated the experiment using Rift Valley fever virus (RVFV) and La Crosse virus (LACV). These bunyaviruses are peculiar in that their transcription depends on concurrent translation (Raju et al., 1989). CHX treatment arrests viral primary transcription at a very early step, but had only a minor influence on IFN-β and ISG56 induction (Figure 1B). Apparently, not even a fully operating primary transcription is required for triggering an IFN response.

Recent reports suggested that particle attachment or membrane fusion can activate an antiviral response (Holm et al., 2012; Noyce et al., 2011), although other groups did not observe this in their systems (Handke et al., 2009; Spirig et al., 2007; Stoltz and Klingström, 2010). We employed several methods to study the involvement of particle attachment. First, we pretreated cells with NH4Cl, an agent that inhibits bunyaviral entry into the cytoplasm (Filone et al., 2006) but does not impede the IFN response (Figure S2A). As shown in Figure 1C, NH4Cl almost entirely abrogated the host response to RVFV. Similarly, three other bunyavirus entry inhibitors (de Boer et al., 2012) (Figure S2B–S2D), as well as virus inactivation by β-propiolactone
primary transcription, but not of replication (Habjan et al., 2009a). tc-VLPs triggered IFN-β and ISG56 expression, even under CHX (Figure 1D). Ghost VLPs, however, did not elicit any IFN response, although they bind equally well to the cells (Figure S2H). These data indicate that, in our system, virus entry or primary transcription is necessary and sufficient for IFN induction.

**The Immediate Early IFN Response Requires Viral 5’Triphosphate RNA and RIG-I**

All viruses used so far contained the RIG-I-activating 5’ppp group on their RNA genome (Habjan et al., 2008a; Hornung et al., 2006; Pichlmair et al., 2006). To clarify the contribution of the genome end, we employed two RNA viruses that lack this feature (see Table S1). Prospect Hill virus (PHV; family Bunyaviridae) has a monophosphate at the 5’ end (Garcin et al., 1995; Habjan et al., 2008a). Semliki Forest virus (SFV; family Togaviridae) has a 5’ cap structure (McInerney et al., 2005). CHX inhibits PHV in the same way as the related RVFV or LACV (Figure S1). SFV is a positive-strand RNA virus (i.e., its genome is directly translated after entry). In the absence of CHX, PHV minimally activated the IFN-β promoter and moderately activated ISG56 transcription, as expected (Prescott et al., 2005), whereas the IFN response to SFV was similar to LACV (Figure 2A). Strikingly, application of CHX completely abrogated the IFN response to both PHV and SFV. In line with this, the transcription factor IRF-3 was activated under CHX-restricted infection with VSV or LACV, but not PHV or SFV (Figure S3A–S3F). Thus, IFN responses to incoming RNA virus nucleocapsids occur only if the genome contains a 5’ppp.

To pin down the responsible RLR, we evaluated the influence of specific knockdowns. Cells were transfected with small interfering RNAs (siRNAs) for MDA5, RIG-I, or a control (Figure S3G), treated with CHX, and infected with VSV or LACV. Knockdown of RIG-I, but not of MDA5, impaired IFN induction by VSV and LACV nucleocapsids (Figure 2B). To bolster these findings, we utilized mouse embryo fibroblasts (MEFs) deficient in RIG-I or MDA5. Because these cells were extremely sensitive to CHX (data not shown), we employed tc-VLPs as a virus system halted at the stage of primary transcription. MEFs lacking MDA5 displayed similar host responses to tc-VLPs as WT MEFs (Figure 2C). MEFs lacking RIG-I, by contrast, completely lost their ability to respond (Figure 2D). Knockout of MAVS (Seth et al., 2005), the adaptor common to MDA5 and RIG-I, had a similar impact (Figure S3H). Taken together, these data suggest that nucleocapsids of 5’ppp RNA viruses activate an IFN response via the RIG-I-MAVS signaling cascade.

**Activation of RIG-I**

Binding of a target RNA to RIG-I triggers a conformational switch (Saito et al., 2007; Takahasi et al., 2008) and oligomerization (Binder et al., 2011; Saito et al., 2007). The conformational switch is indicated by partial resistance to trypsin digestion. We observed that the full infection cycle of SFV, VSV, and LACV leads to the emergence of trypsin-resistant fragments of RIG-I (Figure 3A). When cells had been pretreated with CHX, VSV and LACV still triggered the conformational switch of RIG-I, whereas SFV infection had no effect (Figure 3B). The oligomerization of RIG-I was assayed by native PAGE. In uninfected cells, only monomers of RIG-I were detected (Figure 3C). Curiously, also in SFV-infected cells only monomeric RIG-I is present (Figure 3D), although RIG-I switches conformation (see Figure 3A). For the 5’ppp RNA viruses VSV and LACV, both the full and the CHX-aborted replication cycle resulted in RIG-I oligomerization (Figures 3E and 3F). The 5’-monophosphorylated PHV, by contrast, only weakly activated the conformational switch (Figure 3G) and not the oligomerization (Figure 3H). Taken together, these results suggest that RIG-I is rapidly and strongly activated by viral nucleocapsids in the absence of genome replication, provided the genome is carrying a 5’ppp group.

**Interaction between RIG-I and Incoming Nucleocapsids**

We wondered whether RIG-I could form a complex with viral 5’ppp RNA nucleocapsids. As an experimental system, we chose the setup closest to the immediate early phase (arrested primary transcription), namely infection with bunyaviruses such as LACV under CHX treatment. The N protein thereby served as a marker for viral nucleocapsids. Using confocal immunofluorescence...
microscopy, we detected the nucleocapsids of incoming LACV particles as individual dots in the cytoplasm (Figure 4A). These LACV N dots colocalize with RIG-I (Figure S4A), as indicated by overlay pictures and intensity profiles of the fluorescence signals. Quantitative analysis revealed that LACV nucleocapsids colocalize with RIG-I in 51% of the cases but with the cytoplasmic control protein glyceraldehyde-3-phosphate dehydrogenase (GAPDH) in 33% of the cases (data not shown). N/RIG-I colocalization was also observed in cells undergoing a full replication cycle (data not shown), in agreement with observations on FLUAV (Onomoto et al., 2012). The incoming LACV nucleocapsids also colocalized with peroxisomes (Figure S4B), cytoplasmic organelles involved in immediate early activation of RIG-I (Dixit et al., 2010). Comimmunoprecipitation demonstrated that RIG-I was capable of binding the incoming LACV nucleocapsids (Figure 4B). The same RIG-I/LACV nucleocapsid interaction was observed when a full replication cycle was allowed, whereas MDAS did not interact under any condition (data not shown). Overexpressed N protein, by contrast, could not be precipitated via RIG-I (Figure S4C). We performed similar experiments with RVFV. Again, incoming nucleocapsids triggered the conformational switch and oligomerization of RIG-I and activated IRF-3 (Figure S4D). Moreover, the nucleocapsids of RVFV colocalized (Figure S4E) and coprecipitated with RIG-I (Figure S4F). Strikingly, incoming nucleocapsids could be forced into a high-molecular-weight RIG-I complex. The ATP analog ADP·AlF₃ traps RIG-I in an RNA-bound closed conformation unable to cycle between the different states (Kowalinski et al., 2011). When CHX-treated and infected cells were incubated with ADP·AlF₃, the oligomers of RIG-I shifted from a smear into a single high-molecular-weight band (Figure 4C, upper panels). Probing of the lysates for the viral N protein revealed a similar shift from an oligomeric state to a high-molecular-weight complex (Figure 4C, lower panels). This was true for LACV as well as for RVFV, indicating a general phenomenon. Together, these findings demonstrate that RIG-I is able to associate with nucleocapsids of 5’ppp RNA viruses directly after entry into the cell, leading to RIG-I activation and innate immune signaling.

Nucleocapsid Binding and Activation of RIG-I Are Independent of Mammalian Cofactors

The RIG-I/nucleocapsid interaction could either be direct or mediated by one of the cellular cofactors of RIG-I (Kato et al., 2011; Kok et al., 2011; Miyashita et al., 2011). To distinguish between these possibilities, we employed an insect cell system.
Drosophila melanogaster cells are a useful tool to reconstitute mammalian signaling complexes in a background-free setting (Yang and Reth, 2012) and are infectable with RVFV (Kortekaas et al., 2011). When human RIG-I was expressed in Drosophila D.mel-2 cells, conformational switching and oligomerization were observed after infection with RVFV (Figures 5A and 5B). RIG-I was also activated in vitro by mixing of lysates from RIG-I-transfected D.mel-2 cells with lysates of infected D.mel-2 cells (Figure 5C). Coimmunoprecipitation experiments demonstrated binding of RIG-I to viral nucleocapsids in the Drosophila system (Figure 5D). Again, this nucleocapsid interaction was independent of whether RIG-I contacted RVFV during authentic virus infection or whether lysates containing either RIG-I or RVFV nucleocapsids were mixed in vitro. Thus, activation and nucleocapsid interaction of RIG-I occur in a direct manner and without the contribution of a mammalian cofactor.

Nucleocapsid-Borne 5’ppp dsRNA Is Necessary and Sufficient for RIG-I Activation

Our experiments with bunyaviruses and CHX implicated that the transcriptional activity of nucleocapsids may not be relevant for RIG-I activation. However, CHX still allows some abortive transcription (Raju et al., 1989). To clarify whether RIG-I requires this residual RNA synthesis, we performed several experiments. First, we depleted the cellular NTP pool with the compounds Brequinar (BRQ; inhibits pyrimidine synthesis), mycophenolic acid (MA; reduces GTP levels), pyrazofurin (PYF; reduces CTP and UTP levels), or cyclopentenylcytosine (CPEC; depletes the CTP pool) (Linke et al., 1996; Qing et al., 2010). Each of these inhibitors affected viral RNA synthesis (Figure S5A). Nonetheless, even when combined with CHX, neither inhibitor diminished RIG-I activation by LACV (Figure 6A). In fact, some of the compounds slightly increased RIG-I activation and enhanced the activation of IRF-3 (Figure 6A). Second, we depleted Drosophila D.Mel-2 cell lysates containing either RIG-I or nucleocapsids from NTPs by dialysis. When these dialyzed lysates were mixed with each other, the conformational switch of RIG-I still occurred and was enhanced by adding back ATP (Figure 6B). Third, we employed nucleocapsids that had been isolated from purified virus particles. Dialyzed RIG-I-containing D.Mel-2 extracts were incubated with nucleocapsids of RVFV or LACV. The nucleocapsids clearly induced the conformational switch of RIG-I in vitro (Figure 6C). We conclude from these experiments that viral transcription is not necessary for triggering RIG-I. As outlined above, the genomic RNA of bunyaviruses forms a panhandle structure that has remarkable similarity with the optimal 5’ppp dsRNA ligand identified by transfection and in vitro binding experiments (Figure S5B). Indeed, destroying the dsRNA structures with RNase III or cleaving the triphosphates with a phosphatase abolished the ability of viral nucleocapsids to activate RIG-I (Figures 6D and 6E). Treatment with the single-stranded RNA-specific RNase A, by contrast, had no effect. Thus, a 5’ppp dsRNA structure is indeed necessary for the activation of RIG-I by nucleocapsids.
A Single Site on the Nucleocapsids Is Contacted by RIG-I
We employed ground state depletion (GSD) microscopy to visualize the RIG-I/LACV nucleocapsid complex at 20 nm resolution. This technique allows the display of individual nucleocapsids in their characteristic pseudocircular shape (Figure 7, green channel), which is caused by the dsRNA panhandle formation between the 5' and 3' genome ends (Obijeski et al., 1976). Strikingly, accumulations of RIG-I are contacting nucleocapsids at a single site, lending the cocomplexes a "diamond ring"-like appearance (Figure 7, red channel). The negative control GAPDH, by contrast, appeared to be more distant and not accumulated at the nucleocapsids (Figure S6). These observations add additional weight to our hypothesis that RIG-I binds the nucleocapsids via the terminal 5'ppp dsRNA panhandle.

DISCUSSION
The aim of our study was to clarify whether RIG-I is capable of recognizing the RNA contained within incoming viral nucleocapsids. Our results indicate that this is indeed the case. RIG-I was rapidly activated by viruses with the prototypical 5'ppp dsRNA panhandle structure, even when viral RNA synthesis was abolished. Moreover, we detected a direct, single-site interaction between RIG-I and nucleocapsids that was dependent on 5'ppp dsRNA. Thus, viral nucleocapsids containing a 5'ppp panhandle represent a pathogen-associated molecular pattern (PAMP) for RIG-I. These findings advance the proximal RIG-I-sensitive step of the viral infection cycle from the late stage of genome replication (Baum et al., 2010; Rehwinkel et al., 2010) to the immediate early step of nucleocapsids entering the cytoplasm.

Erroneous, nonencapsidated replication products appear during infection with FLUAV (Vreede et al., 2004). It is quite likely that such naked replication products (Rehwinkel et al., 2010; Vreede et al., 2004), along with defective interfering RNAs (Baum et al., 2010; Killip et al., 2011; Strahle et al., 2006) and newly formed nucleocapsids (this study), are responsible for RIG-I induction under a full viral multiplication cycle. Previous reports already demonstrated that genome replication is not an absolute requirement for IFN induction (Killip et al., 2012; Marcus and Sekellick, 1980) and that transfected nucleocapsids of the 5'ppp viruses measles and VSV can activate IRF-3 even if they are transcriptionally inactive (tenOever et al., 2002, 2004). None of these studies, however, had addressed the involvement of RIG-I or the requirement for a specific 5' genome end. Our work extends their conclusions by showing that incoming viral nucleocapsids can directly activate RIG-I, thus triggering a rapid innate immune response.

It remains to be shown how RIG-I manages to access the encapsidated viral RNA. Although the dsRNA-panhandle structure is covered by the viral polymerase (Resa-Infante et al., 2011), some cytoplasmic exposure is necessary (e.g., to initiate mRNA transcription). The global presence of RIG-I in the
cytoplasm and its high affinity for 5’ppp dsRNA may allow to rapidly entrap a panhandle, even if it is only briefly exposed. Our observation of an early and substantial formation of RIG-I oligomers in response to incoming nucleocapsids supports this hypothesis.

The standard model of RIG-I activation implies that ligand binding induces a conformational change followed by oligomerization (Kowalinski et al., 2011; Saito et al., 2007; Takahasi et al., 2008). Interestingly, oligomerization seems not to be a strict consequence of the conformational switch. While infection with the negative-stranded 5’ppp-RNA viruses VSV, LACV, and RVFV triggers both the conformational switch and the oligomerization, infection with the positive-strand 5’ppp-RNA viruses SFV triggers only the conformational switch. It is known that innate immune recognition of SFV occurs mainly through TLR3 and TLR7/8 (Schulz et al., 2010). VSV, LACV, and RVFV are mainly recognized by RIG-I (Habjan et al., 2008a; Kato et al., 2006; Verbruggen et al., 2010). Thus, a weak RIG-I trigger like SFV may only cause the conformational switch, whereas stronger RIG-I triggers continue to the subsequent oligomerization.

We have introduced Drosophila cells as a tool to establish that RIG-I activation occurs independent of mammalian cofactors. In mammalian cells, RIG-I is fine-tuned by inhibitors and cofactors. The Drosophila cells, which can coexpress up to 12 different recombinant proteins (Yang and Reth, 2012), hold promise as a system to reconstitute the RIG-I signaling complex for studying the details of its regulation.

The classical studies on PRRs and PAMPs involved the usage of purified RNA ligands. However, as we and others (Baum and Garcia-Sastre, 2011; Kato et al., 2011) have pointed out, viral RNAs in their physiological context are complexed with host cell or viral proteins. Recent reports on the PRRs PKR and TLR3 provide increasing evidence that protein-bound RNA ligands are comparable or even more powerful PRR ligands than naked RNAs (Dauber et al., 2009; Lai et al., 2011). It would be interesting to test other PRRs for activation by physiological nucleic acid–protein complexes.

In summary, our results indicate that RIG-I, the major intracellular PRR for viral pathogens, is capable of recognizing the 5’ppp-dsRNA of viral nucleocapsids. This enables a cytoplasmic response to incoming negative-strand RNA viruses at the earliest possible time point of infection.

**EXPERIMENTAL PROCEDURES**

**Cells, Viruses, Plasmids, and Reagents**

A549, 293T, and MEFs deficient in RIG-I, MDA5 (Kato et al., 2006), or MAVS (Seth et al., 2005) were cultured in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal calf serum (FCS) at 37°C and 5% CO₂. D.Mel-2 cells (GIBCO) were cultured in Spodopan (Pan Biotech) at 28°C with no additional CO₂. VSV, SFV, FLUAV, and RVFV were grown in D.Mel-2 cells expressing D.Mel-2 cells, which can coexpress up to 12 different recombinant proteins (Yang and Reth, 2012), hold promise as
Habjan et al., 2008b) were propagated on Vero cells. Plasmids pI.18_RVFV_N, pI.18_RVFV_M, and pI.18_RVFV_L were diluted in DMEM 5% FCS (mammalian cells) or Spodopan (insect cells). Upregulation of inducible genes is depicted in relation to nonstimulated, noninfected (mock) cells.

siRNA Knockdown
Knockdown of gene expression was achieved by 2-fold reverse transfection of siRNAs. siRNAs (25 nM each; see Supplemental Information) were diluted in 100 μL DMEM, mixed with 3 μl HiPerFect (QIAGEN), incubated for 10 min at room temperature, and dropped onto a 12-well plate. Then, 1.5 x 10^5 cells in DMEM 10% FCS were seeded on top. After 48 hr at 37°C, cells were harvested, counted, and 1.5 x 10^5 cells were again reverse transfected as described.

Activation State of RIG-I
To assay the conformation of RIG-I, cells were lysed in PBS/0.5% Triton X-100 and incubated on ice for 10 min. Then, samples were sonified in a Branson 3200 Ultrasonic cleaner at 4°C for 10 min and centrifuged at 4°C for 10 min at 10,000 x g. An aliquot of 25 μg of total protein in 10 μL PBS was digested for 25 min with 0.2 μg/μL L-1-lysylamido-2-phenylethyl chloromethyl ketone-treated trypsin (Sigma-Aldrich) at 37°C. Reaction was stopped by adding 5x sample buffer (250 mM Tris-HCl [pH 6.8], 10% SDS, 50% glycerol, 25% β-mercaptoethanol [β-ME], 0.5% bromphenol blue) and heating at 95°C for 5 min. Samples were subjected to 12% SDS-PAGE and western blot analysis using mouse monoclonal anti-RIG-I antibody (ALME-1; Enzo Life Sciences) at 1:1,000. Staining of the blot with 0.1% Ponceau S in 5% acetic acid served as a loading control.

To investigate the oligomerization of RIG-I, 50 μg of sonified cell lysate in native loading buffer (50 mM Tris-HCl [pH 6.8], 10% glycerol, 0.1% bromphenol blue) was loaded onto a nondenaturing 8% polyacrylamide gel. Proteins were separated by electrophoresis with 50 mM Tris-NaOH (pH 9.0), 384 mM glycine as anode buffer and 50 mM Tris (pH 8.3), 384 mM glycine, 1% sodium deoxycholate as cathode buffer. Western blot analysis was performed as outlined above.

Immunofluorescence Assays
Cells were grown on coverslips to 30%–50% confluency, infected, and incubated for the indicated time. Cells were fixed with 3% paraformaldehyde, permeabilized with 0.5% Triton X-100 in PBS, and washed three times with PBS. Primary antibodies were diluted in PBS/1% FCS. Primary antibodies were either rabbit polyclonal anti-LACV N (1:1,000) (Blakopi et al., 2007) or rabbit polyclonal anti-RVFV N (1:1,000) (Lorenzo et al., 2008) combined with mouse polyclonal anti-human RIG-I (1:200) (Baum et al., 2010). After 1 hr incubation at room temperature, coverslips were washed three times in PBS then treated with goat anti-rabbit Cy2 and goat anti-mouse Cy3 at a dilution of 1:200. After washing three times in PBS, coverslips were mounted with Fluorsave solution (Calbiochem) and examined using a Leica SP5 confocal microscope.

For the superresolution microscopy, a Leica SR GSD microscope was used. Samples were prepared as described for confocal microscopy except that goat anti-rabbit Alexa Fluor 488 (LACV N) and goat anti-mouse Alexa Fluor 647 (RIG-I) were employed as secondary antibody antibodies. Samples were embedded in freshly prepared 100 mM β-mercaptoethanol in PBS (pH 7.4) directly before imaging.

Coimmunoprecipitation Assays
Cells grown in two T75 flasks were scrapped off in 10 ml PBS, centrifuged at low speed, and the pellets lysed in 1,050 μl RIPA buffer (prepared with DEPC-treated H2O) containing protease inhibitors, incubated on ice for 30 min and centrifuged at 4°C for 10 min at 10,000 x g. Supernatants were transferred to fresh tubes and 10% kept as input control. The remaining 90% of the lysates were subjected to immunoprecipitation (IP) using Dynabeads (Invitrogen). In parallel, 1.5 mg beads per IP were coupled with the appropriate antibodies using the Dynabeads antibody coupling kit (Invitrogen). For the anti-p21 and anti-RIG-I-IPs, beads were coupled with rabbit polyclonal anti-mouse antibody (DAKO); for the anti-LACV N IP, beads were coupled with rabbit polyclonal anti-LACV N (each at a 1:20 dilution per IP); and for

Supplemental Information

the anti-RVFV N IP, beads were coupled with mouse polyclonal anti-N serum (Habjan et al., 2009b) for 20 hr at 37°C. After antibody coupling, beads were washed two times with RIPA buffer, and the anti-mouse antibody beads were further incubated with mouse mononal anti-p21 (Santa Cruz Biotechnology) or mouse monoclonal anti-RIG-I antibody ALME-1 at a 1:200 dilution in RIPA buffer for 2 hr at 4°C. The beads were then incubated with the lysates for 2 hr at 4°C and the immunoprecipitates washed three times with RIPA buffer and then eluted with sample buffer (50 mM Tris-HCl [pH 6.8], 2% SDS, 10% glycerol, 5% β-ME, 0.1% bromphenol blue) for 5 min at 95°C. Eluates were analyzed by western blotting using rabbit polyclonal anti-LACV N or anti-RVFV N antisera (1:11,000), mouse monoclonal anti-RIG-I antibody ALME-1 (1:1,000), or mouse monoclonal anti-p21 (1:500). Protein A horseradish peroxidase conjugate (Millipore; 1:10,000) was used for detection.

Supplemental Information includes six figures, one table, and Supplemental Experimental Procedures and can be found with this article online at http://dx.doi.org/10.1016/j.chom.2013.01.012.

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REFERENCES


Incoming RNA Virus Nucleocapsids Containing a 5′-Triphosphorylated Genome Activate RIG-I and Antiviral Signaling


Table S1

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1Virus abbreviations: FLUAV, influenza A virus; LACV, La Crosse virus; PHV, Prospect Hill virus; RVFV, Rift Valley fever virus; SFV, Semliki Forest virus; VLPs, virus-like particles; VSV, Vesicular stomatitis virus. 2(-): negative-strand RNA virus, (+): positive-strand RNA virus. 3CHX, Cycloheximide. 4n.a., not applicable
Figure S1. Effect of CHX Treatment on Viral RNA Synthesis (related to all figures)

A549 cells were either left untreated, or pretreated with CHX (50 µg/ml) for 1 h at 37°C, and infected with viruses at an MOI of 1. Seven hours (RVFV) or twelve hours (all other viruses) post-infection, cells were lysed and the total cell RNA was extracted. Viral RNA levels were measured by real-time RT-PCR as indicated in Supplemental Experimental Procedures.
Figure S2

(A) IFN-beta mRNA

(B) ISG56 mRNA

(C) IFN-beta mRNA

(D) ISG56 mRNA

(E) IFN-beta mRNA

(F) ISG56 mRNA

(G) Western Blot

(H) Immunofluorescence
Figure S2. Control Experiments, Methods, and Tools for Investigating IFN Induction by Incoming Virus Particles (related to Figure 1)

(A) ISG response to IFN is unaffected by NH4Cl. A549 cells were treated for 1 h with 0 or 50 mM NH4Cl, and then either transfected with 500 ng RNA of RVFVΔNSs::GFP particles (vRNA), or treated with 1000 U/ml IFN-α (IFN). IFN-β and ISG56 mRNA levels were analyzed 24 h later by real-time RT-PCR. (B, C, and D) RVFV entry inhibitors and the IFN response. (B) A549 cells were treated for 1 h either with 8 µl DMSO, or with 100 nM of Bafilomycin A (BafA) in DMSO, and infected and analysed as described for Figure 1C. (C) A parallel experiment employing 160 µM of Dynasore (DynS), or 40 µM of Chlorpromazine (ChPZ), each dissolved in DMSO. (D) Real-time RT-PCR analysis for the RVFV L gene as described (Bird et al., 2007). (E and F) IFN response to inactivated virus particles. A549 cells were infected with RVFVΔNSs::GFP particles inactivated with β-propiolactone (β-PL) (E) or irradiated with UV (λ=311 nm) for the indicated period of time (F). (G) Production of RVFV VLPs. (Upper left panel) Biological activity of GFP-expressing tc-VLPs. BSR-T7/5 cells were transfected with plasmids for RVFV N and L and 24 h later infected with tc-VLPs. At 24 h post infection, GFP-positive cells were observed by fluorescence microscopy. (Upper right panel) Concentrated tc-VLPs or ghost VLPs, or cell lysate of RVFV-infected Vero cells were loaded on a 12% polyacrylamide gel and tested by immunoblot using pan-specific anti-RVFV mouse antiserum. (Lower panel) Ghost VLP preparations contain intact particles which protect the N from trypsin digestion. Recombinant RVFV nucleoprotein rN (0.2µg) containing a His-tag (His-N), as well as RVFVΔNSs::GFP, tc-VLPs, and ghost VLPs purified by ultracentrifugation were exposed to trypsin (100 µg/ml), and incubated for 30 min at 37°C. Samples were immunoblotted using an anti-RVFV N antibody. Fractions of trypsin-sensitive N indicate the presence of free N secreted by infected and transfected cells, as observed earlier (Kortekaas et al., 2011). (H) Attachment of VLPs. A549 cells were incubated with RVFV VLPs that are either complete (tc-VLPs), or for which either the polymerase (-L), the glycoproteins (-M), or the polymerase and the minireplicon (ghost VLPs) were omitted. RVFVΔNSs::REN infection served as positive control (RVFV). Cells were infected for 1 h at 4°C, fixed with 3% paraformaldehyde and analyzed by immunofluorescence using rabbit anti-RVFV N antisera (1:200, green channel) and Hoechst 33342 (blue channel) as nuclear counterstain. Note that cells were not permeabilized prior to analysis. Therefore only particles attached to the cells are detected. Figures S2A to F show mean values and standard deviations from three independent experiments.
Figure S3. Companion Experiments and Controls for the Influence of the 5’ppp Group and RIG-I / MAVS Signaling on IFN Induction (related to Figure 2)

(A to F) Activation of IRF-3 in dependence of viral 5’ genome ends. A549 cells were left untreated (UT) or treated with 50 µg/ml CHX. Then, cells were either mock infected (A) or infected with SFV (B), VSV (C), or LACVdelNSs (D) as described for Figure 1. At the indicated time points post-infection, cells were lysed with RIPA buffer and protein extracts cleared from cell debris separated by native PAGE. The presence of phospho-IRF-3 (P-IRF-3) and actin, and the oligomerization state of total IRF-3 were analyzed by Western blot using specific antisera. As internal controls, lysate from VSV-infected cells at 6 h post-infection was used as positive control for the mock infected cells (CTRL), and lysates from mock infected cells were used as negative control for the virus-infected cells. (E) Subcellular localization of IRF-3 in LACV-infected cells. A549 cells were treated with CHX or left untreated, and then mock infected or infected with LACVdelNSs. Upper panel shows representative examples of immunofluorescence analyses for IRF-3 (green). Graph shows statistics (mean and standard deviations) of the IRF-3 subcellular localization from a total of 4 independent experiments, in each of which 100 cells were randomly chosen. (F) PHV and IRF-3 activation. A549 cells were mock infected or infected with LACVdelNSs or PHV, and 6 h later monitored for IRF-3 phosphorylation and dimerisation as described for panels A to D. (G) Knockdown of intracellular PRRs. A549 cells were transfected with a negative control siRNA (CTRL), or with siRNAs directed against MDA5 or RIG-I. After two rounds of transfection, cells were either infected as indicated for Figure 2B, or lysed and tested for the presence of the siRNA targets by western blot analysis. MDA5 was immunodetected using rabbit polyclonal antiserum NBP1-03299 (1:500; Novus Biologicals), actin and RIG-I were immunodetected as indicated for A–D and Figure 3, respectively. (H) Host response to RVFV primary transcription in MAVS knockout cells. wt MEFs or MEFs lacking the MAVS gene were infected with transcriptionally competent RVFV VLPs expressing a GFP reporter minireplicon (tc-VLPs). After 24 h of infection, mRNA levels of IFN-β and ISG56 were determined by real-time RT-PCR. Mean values and standard deviations from three independent experiments are shown.
Figure S4. Companion Experiments and Controls for the Interaction of RIG-I with Viral Nucleocapsids (related to Figure 4)

(A) RIG-I signal in knock-down cells. A549 cells were transfected with a negative control siRNA (CTRL), or with an siRNA against RIG-I. After two rounds of transfection, cells were tested for the presence of RIG-I by Western blot (upper panel) and immunofluorescence analysis (lower panel). (B) Co-localization of LACV nucleocapsids with peroxisomes. CHX-treated A549 cells were infected with LACVdelNSs (MOI 5) and 5 h later analyzed by immunofluorescence using rabbit and mouse antisera directed against LACV N (green channel) or catalase as a marker for peroxisomes (red channel), respectively. Cell nuclei were counterstained with DAPI (blue channel). The square area of the inset is digitally magnified on the right hand side. Three fluorescence intensity profiles (numbered 1 to 3) are shown on the bottom. (C) Only viral nucleocapsids, but not the nucleocapsid protein alone, interact with RIG-I. A549 cells were either untreated, transfected for 24 h with plasmid pI.18-LACV N, or infected for 5 h with LACVdelNSs (MOI of 5). Cells were lysed in RIPA buffer and subjected to immunoprecipitation (IP) for LACV N or RIG-I and analyzed by Western blot. As input control, 10% of the cell lysate were analyzed in parallel (left lanes). * indicate unspecific bands. (D) Activation of RIG-I and IRF-3 by RVFV. (Upper panel) A549 cells were treated with 0 or 50 µg/ml CHX and then infected with RVFVΔNSs::REN at an MOI of 5. After the indicated incubation periods, cells were lysed in PBS / 0.5% Triton X-100 and analysed for oligomerization of RIG-I and IRF-3, and for phosphorylation of IRF-3. (Lower panel) RIG-I conformational switch assay of A549 cells treated with CHX and infected for 5 h with RVFVΔNSs::REN. (E) Co-localization of RIG-I with RVFV nucleocapsids. CHX-treated A549 cells were infected with RVFVΔNSs::GFP (MOI 5) for 5 h and analyzed by immunofluorescence using a rabbit antiserum against RVFV N (1:5000; green channel) and mouse polyclonal antiserum against RIG-I (1:200; red channel), respectively. Cell nuclei were counterstained with DAPI (blue channel). The square area of the inset is digitally magnified on the right hand side. Three fluorescence intensity profiles (numbered 1 to 3) are shown on the bottom. (F) Co-immunoprecipitation of RIG-I and RVFV nucleocapsids. CHX-treated A549 cells were infected with RVFVΔNSs::GFP (MOI 5) and 5 h later lysed in RIPA buffer. Lysates were subjected to immunoprecipitation (IP) using rabbit antiserum against RVFV N (1:1000), mouse monoclonal anti-RIG-I antibody ALME-1 (1:1000), or mouse monoclonal anti-p21 (1:500). As input control, 10% of the cell lysate were analyzed in parallel (left lanes).
Figure S5. Control Experiment and Illustration for the Activation of RIG-I by the Viral Panhandle (related to Figure 6).

(A) Effect of NTP withdrawal on LACV RNA synthesis. A549 cells pretreated with different inhibitors were infected with LACVdelNSs for 5 h at an MOI of 5. Before and during the infection period, the cell culture medium contained either no additive (-), CHX (50 µg/ml) added 1 h before infection, or BRQ (10 µM, stocks dissolved in DMSO), MA (10 µM, stocks dissolved in methanol), PYF (10 µM, stocks dissolved in DMSO), or CPEC (5 µM, stocks dissolved in DMSO) added 24 h before infection. Total cellular RNA was isolated and analyzed by real-time RT-PCR using primers specific for the LACV N RNA (Verbruggen et al., 2011).

(B) LACV panhandle structure. The first 32 nucleotides of the 5’ and 3’ LACV M genome segment (Genbank acc # NC_004109) are depicted as an example. Note that the non-Watson–Crick base pair G-U (bold face characters) does not disturb the dsRNA structure (Holbrook et al., 1991).
**Figure S6.** Super-Resolution Immunofluorescence Microscopy of GAPDH and LACV Nucleocapsids (related to Figure 7). A549 cells were treated with 50 µg/ml CHX and infected with LACVdelNSs (MOI 5). Cells were analyzed 5 h later by GSD double immunofluorescence using antisera against LACV N (green channel) or glyceraldehyde-3-phosphate dehydrogenase (GAPDH, red channel). The primary antibody anti-GAPDH mouse monoclonal 9B3 (HyTest) was diluted 1:100. Four example areas with nucleocapsids are shown. Scale bar 200 nm.
Supplemental Experimental Procedures

**Primers for real-time RT-PCR**
mRNAs of human IFN-β, ISG56, and γ-actin were measured with QuantiTect primers QT00203763, QT00201012, and QT00996415, respectively. mRNAs of murine IFN-β, ISG56, and GAPDH were measured with QuantiTect primers QT00249662, QT01161286, and QT01658692, respectively.

**Short interfering RNAs**
AllStar Negative Control siRNA and validated FlexiTube siRNAs (Qiagen) against RIG-I mRNA (SI03649037) and MDA5 mRNA (GS23586) were used.

**Chemical inactivation of virus particles**
Virus stocks containing 0.05% β-propiolactone (Acros Organics) were incubated at 4°C for 16 h. β-propiolactone was subsequently hydrolyzed by incubating samples at 37°C for 2 h.

**Homodimerisation and phosphorylation of IRF-3**
A549 cells were grown to 80% confluency in T25 flasks. Cell lysis and native-polyacrylamide gel electrophoresis (PAGE) were performed as described (Iwamura et al., 2001), with minor modifications. Briefly, cells were lysed in 0.5% Triton X-100 dissolved in PBS supplemented with protease inhibitor cocktail (Roche) and incubated on ice for 10 min. Lysates were sonified in a Branson 3200 Ultrasonic cleaner for 10 min at 4°C and then centrifuged at 4°C for 10 min at 10,000×g. Protein concentration was determined by Bradford assay (BioRad) and 50 µg of total protein lysate were separated by electrophoresis in a nondenaturing 8% polyacrylamide gel, with 1% sodiumdeoxycholate in the cathode buffer. Total IRF-3 and phosphorylated IRF-3 monomers and oligomers were detected by Western blot analysis with rabbit polyclonal anti-IRF-3 antibody FL-425 (1:500, Santa Cruz Biotechnology) or rabbit monoclonal anti-P386-IRF-3 (1:100, IBL), respectively. β-actin, detected by mouse monoclonal antibody 8H10D10 (1:1000, Cell signaling) served as a loading control.

**Subcellular localization of IRF-3**
Cells were analyzed by immunofluorescence (see Experimental Procedures) using the FL-425 rabbit polyclonal anti-IRF-3 as primary antibody (1:500, Santa Cruz Biotechnology). And goat anti-rabbit Cy2 conjugated as secondary antibody.
Real-time RT-PCR for detection of viral sequences

Total cellular RNA was isolated with the NucleoSpin RNA II kit (Macherey-Nagel) and eluted in 30 µl of ddH2O. An aliquot of 600 ng RNA was then used as a template for cDNA synthesis. For detection of FLUAV and VSV sequences, reverse transcription was performed using the Qiagen QuantiTect Reverse Transcription Kit using the (-) strand specific forward primers 5’-GAT AGT ACC GGA GGA TTG ACG ACT A-3’ (VSV) and 5’-GGA CTG CAG CGT AGA CGC TT-3’ (FLUAV). PCR was performed using QuantiTect SYBR Green PCR Kit and the respective forward primers, combined with the reverse primers 5’- TCA AAC CAT CCG AGC CAT TC-3’ (VSV) and 5’- CAT CCT GTT GTA TAT GAG GCC CAT-3’ (FLUAV). For detection of LACV, PHV and SFV sequences, the one-step QuantiTect SYBR Green RT-PCR Kit was used. Primer sequences were 5’-GGG TAT ATG GAC TTC TGT G-3’ and 5’-GCC TTC CTC TCT GTC TTA-3’ (LACV S segment forward and reverse (Verbruggen et al., 2011)), 5’-CTC AAA ATT GGC AGC TAC A-3’ and 5’-CTT CAC CGG CAG GCT G-3’ (PHV S segment forward and reverse, and 5’-GCA AGA GGC AAA CGA ACA GA-3’ and 5’-GGG AAA AGA TGA GCA AAC CA-3’ (SFV nsp3 forward and reverse (Fragkoudis et al., 2008)). mRNA levels of human γ-actin were detected with QuantiTect primers QT00996415. All virus-specific values were normalized against the γ-actin mRNA signal using the ddCT method. Up-regulation of viral sequences is depicted in relation to non-stimulated, non-infected (mock) cells.
Supplemental References


4.3. Influenza virus adaptation PB2-627K modulates nucleocapsid recognition by the pathogen sensor RIG-I

Own contribution:

I performed or contributed to all experiments. Work in biosafety level 3 was performed by Hanna Sediri (group of Prof. Dr. Klenk, Institute for Virology, Marburg) while I performed analysis of the samples. I contributed to writing of the manuscript.

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Influenza virus adaptation PB2-627K modulates nucleocapsid inhibition by the pathogen sensor RIG-I

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Running title: Influenza virus PB2-627K counteracts RIG-I  
Key words: influenza virus; nucleocapsids; 5' triphosphate dsRNA; panhandle; RIG-I; signaling-independent virus inhibition; PB2-627K; PB2-627E
Summary

The cytoplasmic RNA helicase RIG-I plays a key role in innate immune-sensing of RNA viruses and antiviral signaling. We show that RIG-I recognizes incoming nucleocapsids of influenza A virus (FLUAV) during their short cytoplasmic transit to the nucleus. RIG-I activation was mediated by 5'-triphosphorylated dsRNA on nucleocapsids and modulated by polymorphisms at position 627 of polymerase subunit PB2. Avian FLUAV nucleocapsids possessing PB2-627E were prone to an increased RIG-I recognition as compared to those with mammalian-adapted PB2-627K. Previously known phenotypes of PB2-627E in mammals such as delayed onset of infection and reduced nucleocapsid stability were partially restored in RIG-I-defective cells. The inhibitory effect of RIG-I on PB2-627E nucleocapsids was independent of antiviral signaling. These results identify RIG-I as a directly acting mammalian restriction factor for avian FLUAV and highlight nucleocapsid disruption as a promising target for antiviral therapy.
Highlights

► 5’ppp dsRNA panhandle of incoming influenza virus nucleocapsids activates RIG-I
► Human-adaptive mutation PB2-627K counteracts activation of RIG-I
► RIG-I directly inhibits incoming nucleocapsids with the avian PB2-627E signature
► Strength of polymerase binding to NP determines the sensitivity to RIG-I
Introduction

Influenza A viruses (FLUAV; family Orthomyxoviridae) are a significant health threat. Regular, global outbreaks are occurring due to replenishment from a seemingly unlimited reservoir of strains in birds (Cauldwell et al., 2014; Klenk, 2014; Manz et al., 2013). FLUAV virions consist of a lipid envelope with the glycoproteins HA and NA, that also provide the basis for taxonomic classification, as well as the proton channel M2. The inner leaflet of the viral envelope is covered by the M1 protein. Inside the particles are eight nucleocapsids containing the negative-strand RNA genome encapsidated by the nucleoprotein (NP) and associated with the RNA polymerase complex with subunits PB1, PB2, and PA. The eight genome segments encode the main structural proteins, auxiliary proteins (NS1, NEP), and sometimes other, strain-dependent, factors. Gene expression is regulated by partially complementary 5′ and 3′ end sequences (the “panhandle”) that pseudo-circularize the single-stranded RNA genome and serve as promoter for transcription and replication. An outstanding feature of FLUAV is that most parts of the replication cycle occur in the nucleus.

RIG-I (Retinoic acid-inducible gene I) is an RNA helicase that acts as a major host sensor for virus infections in the cytoplasm (Yoo et al., 2014). RIG-I recognizes non-self RNA structures and activates a signaling cascade leading to phosphorylation of transcription factor IRF3 and induction of the antiviral type I interferons (IFN-α/β).

RIG-I contains a central multipartite helicase/ATPase domain flanked by N-terminal caspase recruitment domains (CARDs) and a C-terminal domain (CTD) (Yoo et al., 2014). Unstimulated RIG-I is in an auto-repressed, monomeric conformation with the CARDs confined and the CTD exposed. When the CTD binds a ligand RNA, RIG-I undergoes a conformational switch enabling it expose the CARDs for interaction with TRIM25 and MAVS to initiate antiviral signaling (Gack, 2014; Kowalinski et al., 2011; Rawling and Pyle, 2014).
Synthetic short 5’ppp-dsRNA represents the optimal RIG-I ligand (Schlee et al., 2009; Schmidt et al., 2009), a structure with remarkable similarity to the panhandle formed by the annealed 5’ and 3’ genome ends of segmented negative-strand RNA viruses (Schlee, 2013; Weber and Weber, 2014b). Accordingly, naked genomic RNA of FLUAV and other negative-strand RNA viruses is an excellent activator of RIG-I (Habjan et al., 2008; Hornung et al., 2006; Pichlmair et al., 2006). We recently reported that RIG-I is capable of recognizing 5’ppp-dsRNA panhandles also in the physiological context, when packaged by viral nucleoprotein and polymerase into nucleocapsids (Weber et al., 2013). That study, however, had focused on the cytoplasmically replicating bunyaviruses. Therefore, it remained open whether the panhandles of FLUAV nucleocapsids could also be sensed by the cytoplasmic RIG-I, and whether this occurs during the passage of the incoming nucleocapsids from the endosome to the nucleus early in infection. Here, we report that the transiting FLUAV nucleocapsids are indeed recognized and directly impaired by RIG-I, and that the degree of RIG-I sensitivity varies depending on adaptive mutations in the polymerase subunit PB2.
Results

Activation of RIG-I by incoming FLUAV nucleocapsids

At the onset of infection, the nucleocapsids of FLUAV are released from endosomes to be transported through the cytoplasm and into the nucleus. Only there they start primary transcription, followed by translation and genome replication. To investigate the interaction between incoming nucleocapsids (and not their RNA products arising later on) and RIG-I in the cytoplasm, we (i) synchronized infection by incubating the virus inoculum at 4°C for 1 h, (ii) allowed the subsequent infection at 37°C to proceed for only 1 h, and (iii) added various inhibitors to ensure restriction to the immediate early infection phase. The inhibitors were Cycloheximide (CHX; blocks protein synthesis and therefore viral genome replication), leptomycin B (LMB; inhibits nuclear export of nucleocapsids), and actinomycin D (ActD; inhibits viral transcription) (Figure S1A). In addition, we tested Ivermectin (IVM) which is known to block the nuclear import machinery (Wagstaff et al., 2012). Human A549 cells were pretreated with one or several of the inhibitors, and 1 h-infected with strain A/PR8/34 (H1N1) at a multiplicity of infection (MOI) of 1 as outlined above. Irrespective of the inhibitor, incoming FLUAV nucleocapsids triggered two markers of RIG-I activation: the formation of homo-oligomers (Figure 1A), and the conformational switch reflected by a partial trypsin resistance (Figure 1B). RIG-I activation by incoming nucleocapsids was comparable to levels obtained by 1 h-transfection of naked virus particle RNA (vRNA). Quantification by RT-qPCR demonstrated the expected absence of viral RNA synthesis (Figures 1C and S1B). In line with this, transcription inhibition by various NTP depleting agents also had no influence on RIG-I activation by FLUAV (Figure S1C). Nonetheless, not only RIG-I, but also its downstream target IRF3 were activated by the incoming nucleocapsids (Figure 1D). It must be remarked that in all our experiments, virus stocks were used which were free of defective interfering particles (data not shown), a known activator of RIG-I (Baum et al., 2010).
Moreover, in agreement with a previous study (Killip et al., 2014) we noted that addition of ActD to the mix containing CHX and LMB results in occasional background activation of antiviral signaling in uninfected cells (data not shown). We therefore conducted the majority of subsequent experiments with CHX/LMB treatment and within the short, 1 h-period of synchronized infection, conditions that were optimal for robustly studying interactions of RIG-I with incoming nucleocapsids.

**RIG-I binds to the panhandle on FLUAV nucleocapsids**

Confocal microscopy revealed that the incoming nucleocapsids are co-localizing with RIG-I, but not with the related helicase MDA5 (Figure S2A and data not shown). Superresolution microscopy suggests that RIG-I directly attaches on one side of the rod-like nucleocapsids (Figure 2A). The stability of these co-complexes was tested by pull-down experiments. Although the viral input was barely detectable as expected for a 1 h infection, the RIG-I immunoprecipitates contained enriched amounts of NP (Figure 2B), indicating interaction of RIG-I with incoming nucleocapsids. Again, this was not dependent on viral RNA synthesis (Figure S2B). When cell lysates were separated in a CsCl gradient, we observed a partial, virus-induced shift of RIG-I from higher density fractions towards the lower density fractions which also contained the nucleocapsids (Figure 2C; fractions 7,8,9 in uninfected cells vs fractions 3,4 and 6 in infected cells). RIG-I was also activated and shifted towards lower density CsCl fractions when its ATPase activity was inhibited with ADP•AlF₃, suggesting that complex formation is independent of downstream events (Figures S2C-S2E). We also used our insect cell / *in vitro* system (Weber et al., 2013). Extracts of *Drosophila* S2 cells expressing human RIG-I were dialyzed and supplemented with ATP (to support RIG-I activation). The recombinant RIG-I reacted to purified and dialyzed FLUAV nucleocapsids by conformational switching, oligomerization, and a shift of RIG-I fractions in the CsCl gradient (Figure S2F-S2H). To test the structural determinants of RIG-I activation, purified
FLUAV nucleocapsids were pretreated in vitro with enzymes. Both destruction of dsRNA by RNase III as well as cleavage of the 5’ppp by a phosphatase aborted RIG-I stimulation, whereas the ssRNA-specific RNase A had no such effect (Figure 2D). Importantly, RIG-I activation did not depend on the specific nucleocapsid preparation method, and was also observed for nucleocapsids that were affinity-purified via a Strep-tagged PB2 subunit (Figure S2I-S2K). Also cotransfection experiments demonstrated that the pull-down of NP by RIG-I is dependent on the genomic RNA, and not on protein-protein interactions (see below). Together, these data suggest that RIG-I directly interacts with the 5’ppp dsRNA panhandle on intact FLUAV nucleocapsids and in a manner that is independent of mammalian cofactors or viral RNA synthesis.

PB2 is a RIG-I antagonist

Avian FLUAV strains need to acquire adaptive mutations to establish infection in mammals. A major determinant of host switching and virulence is the polymerase subunit PB2 (Hatta et al., 2001). PB2 position 627, in particular, carries in avian strains a glutamic acid (E), but in most mammalian-adapted strains a lysine (K) (Subbarao et al., 1993). The reason for the mammalian selection pressure towards PB2-627K is not fully understood (Cauldwell et al., 2014; Manz et al., 2013). Interestingly, however, chicken are known to be deficient in RIG-I (Barber et al., 2010). Using the conformational switch assay, we investigated whether RIG-I might be involved in the mammalian-specific effects on avian-signature PB2 polymerases. Human A549 cells were exposed to the immediate early infection phase of variants of four FLUAV strains, A/quail/Shantou/2061/00 (H9N2), A/Thai/KAN-1/04 (H5N1), pandemic A/Hamburg/05/2009 (pH1N1), or A/WSN/33 (H1N1). In all cases, those viruses with the avian signature PB2-627E activated RIG-I much stronger than those with the mammalian signature PB2-627K (Figure 3A and S3A). These differences were not due to variations in input RNA or RNA synthesis, as viral RNA levels were comparable and did not increase
during the 1 h-experiment (Figure 3B and S3B-S3D). Also in CsCl gradient assays, we observed a more pronounced shift of RIG-I fractions in response to a PB2-627E virus (Fig. 3C, left panels). The PB2-627E virus also relocalized the RIG-I interactors MAVS and TRIM25 (Fig. 3C, right panels), further supporting the notion of a stronger RIG-I activation by the avian-signature nucleocapsids.

The A/PR/8/34 strain used for the initial RIG-I activation experiments (see Figures 1 and 2) contains PB2-627K (Foeglein et al., 2011). A comparison of A/PR/8/34 with PB2-627K and -627E variants of A/WSN/33 (H1N1) demonstrates its relatively weak RIG-I activation potential (Figure S3E), thus being in line with the correlation between reduced RIG-I activation and the PB2-627K signature. Virus-like particles containing a A/WSN/33 (H1N1) reporter minigenome (VLPs) showed the same PB2-627-dependent phenotype, independent of the particular envelope protein (Figure S3F). This confirms that nucleocapsids are the critical component. As expected, RIG-I activation by PB2-627E virus was independent of any viral RNA synthesis (Figure S3G). We also measured IFN induction obtained after overnight infection by the different FLUAV strains, again under CHX and LMB. Surprisingly, despite the clear effects on RIG-I activation described above, there were no consistent PB2-627-dependent differences in IFN induction, both in wt and in RIG-I- depleted knockdown cells (Figure 3D and S3H). Of note, human-adapted PB2-627K has a higher polymerase activity in mammalian cells (Naffakh et al., 2000), but nonetheless activated RIG-I much weaker than PB2-627E. This again argues against RNA synthesis as a trigger of RIG-I. Collectively, our data indicate that the adaptive mutation PB2-627K represents a viral countermechanism to RIG-I recognition, even if it has no significant effect on IFN induction.
RIG-I influences infection by incoming PB2-627E nucleocapsids

PB2-627E confers reduced polymerase activity in mammalian cells, whereas in avian cells the activity is similar to PB2-627K (Massin et al., 2001; Paterson et al., 2014). The species-specific differences were attributed to a mammalian restriction factor acting on PB2-627E only (Mehle and Doudna, 2008). We wondered whether RIG-I could be contributing to PB2-627E suppression in mammalian cells. The classic assay for measuring FLUAV polymerase activity is the minireplicon reporter system, consisting of plasmid-expressed polymerase subunits and NP that encapsidate, transcribe and replicate a reporter minigenome bearing panhandle sequences. Minireplicon activity in human 293 wt cells confirmed the difference between the two PB2 variants (Figure 4A). We generated knockout 293 cells (Figure S4A) to test the effect of RIG-I on minireplicon activity, but could not detect major differences to wt cells or MDA5 knockout cells (see Figure 4A). It needs to be noted, however, that here the recombinant nucleocapsids are not entering the cytoplasm like in infection, but are artificially assembled in the nucleus by the plasmid-encoded proteins. Thus, as an alternative closer to the immediate early infection situation, we employed VLPs containing recombinant FLUAV nucleocapsids. Reporter activity in the 293 wt cells used to produce VLPs for strain A/WSN/33 (H1N1) differed between the two PB2 variants, as expected (Figure 4B and S4B). However, VLPs with PB2-627E re-gained considerable activity upon infection of ΔRIG-I cells, but not in wt cells or ΔMDA5 cells (Figure 4C). Curiously, in cells that lacked the RIG-I adaptor MAVS, PB2-627E could also not be rescued, suggesting that signaling may not be necessary (see below). We also tested the influence of RIG-I on viral multiplication. To obtain multicycle growth, we infected 293 wt or ΔRIG-I cells for 24 h with MOI 0.0001 of PB2 variants of strain A/Thai/KAN-1/04 (H5N1). The absence of RIG-I rescued the low yields of the PB2-627E virus by one order of magnitude, whereas the PB2-627K virus was slightly reduced (Figure 4D). Overall, the approximately 50,000 fold growth difference between the PB2 signature viruses in wt cells decreased to 400 fold in ΔRIG-I cells. Since
RIG-I is activated by nucleocapsid entry, we also monitored the initial establishment of infection. Wt or ΔRIG-I cells were infected with MOI 1, and the synthesis of NP assayed for 8 h. For PB2-627K viruses, NP was detected from 4 h p.i. on in both cell types (Figures 4E, S4C and S4D). For the PB2-627E virus, by contrast, NP was detected at 8 h p.i. in wt cells, but from 6 h on in ΔRIG-I cells. Since viral entry is neither affected by the PB2 signature nor by the cell genotype (Figure S4E), we conclude that RIG-I targets PB2-627E-type nucleocapsids during the onset of infection.

Species- and signaling-independent antiviral effect of RIG-I

An unexpected result of our VLP infection experiments (see figure 4C) was that the attenuated phenotype of incoming PB2-627E nucleocapsids was not rescued by deleting the signaling adaptor MAVS. Similarly, also virus with PB2-627E was delayed in ΔMAVS cells (Figure 5A). To investigate this further, we transcomplemented ΔRIG-I cells with an ATPase-negative RIG-I mutant (K270A) that is unable to signal but still able to bind RNA (Takahashi et al., 2008). Strikingly, the RIG-I K270A mutant was as potent as wt RIG-I in delaying the onset of PB2-627E virus and VLP infection (Figures 5B and S5). This is in line with our observation that chemical ATPase inhibition of RIG-I does not impede activation and nucleocapsid binding (see Figures S2C to S2E). Transcomplementation with human RIG-I also endowed chicken cells with the ability to slow down PB2-627E virus infection (Figure 5C). Again, this was independent of antiviral signaling. These data suggest that the binding of RIG-I to incoming nucleocapsids (see Figs. 2 and S2) is sufficient to delay infection by FLUAV strains with the avian PB2-627E signature. Thus, RIG-I signaling may not be required for the restriction of avian strains early in infection.

Nucleocapsid stability influences RIG-I binding and sensitivity
The 627K adaptation is known to increase the binding of PB2 to NP (Labadie et al., 2007).

Although this had been disputed as an artefact of the stronger polymerase activity (and hence nucleocapsid formation) by PB2-627K (Cauldwell et al., 2013), others had shown stronger NP binding by PB2-627K also in a nucleocapsid-free in vitro system (Ng et al., 2012). In our immediate-early infection setup, infection occurred for 1 h and the formation of daughter nucleocapsids was impossible due to CHX treatment. Nonetheless, also under these conditions NP co-precipitated more PB2-627K than PB2-627E, confirming the adaptation-specific differences in binding affinity (Figure 6A). Moreover, NP / PB2-627E interactions slightly increased in ΔRIG-I cells that were infected (see Figure 6A) or are harbouring minireplicon systems (Figures S6A and S6B).

RIG-I directly interacts with the panhandles of FLUAV nucleocapsids (see Figures 2D and S2K). Interestingly, RIG-I pulled down more incoming nucleocapsids of the PB2-627E type than of the PB2-627K type (Figure 6B). Moreover, the PB2 protein of the 627K type was found to be co-precipitated, but not PB2 of the 627E type. Similar results were obtained with recombinant nucleocapsids of the minireplicon system, an experimental set-up that also allowed to demonstrate that RIG-I-NP interactions are depending on the encapsidated genome RNA (Figure 6C). Thus, nucleocapsids of the PB2-627E type are more efficiently bound by RIG-I.

The lower affinity of PB2-627E to NP may enable RIG-I to better access the nucleocapsid-borne panhandle RNA. This would implicate that the strength of the polymerase - nucleocapsid interaction influences RIG-I activity. To directly test this hypothesis, we disrupted the polymerase complex with a PB1-derived peptide (PB1-T6Y) (Wunderlich et al., 2011). The inhibitor peptide massively increased RIG-I activation by incoming nucleocapsids, and independent of the PB2-627 signature (Figures 6D and S6C). Thus, the strength of polymerase binding, modulated either naturally (by PB2 mutation) or artificially (by disrupting PB1-PA interactions), sensitizes nucleocapsids to RIG-I.
In summary, our results indicate that FLUAV nucleocapsids are prone to signaling-independent RIG-I inhibition during their passage through the cytoplasm early in infection. RIG-I recognizes the panhandle structure on the viral genome RNA, which is normally bound by the polymerase complex. The sensitivity to RIG-I is determined by the well known host adaptation at PB2-627, which affects the binding affinity of PB2 to nucleocapsids. Thus, RIG-I apparently represents one of the mammalian restriction factors driving adaption of avian FLUAV strains towards tighter polymerase binding.
Discussion

An outstanding feature of influenza viruses is the replication in the nucleus. This allows access to the cellular transcription and splicing machineries, and - as our results suggest - the hiding from cytoplasmic RIG-I during most phases of the infection cycle. In fact, since orthomyxoviruses are also unique with respect to the unusually high number of genome segments (and hence RIG-I ligands), it is possible that moving into the nucleus initially evolved as a RIG-I counterstrategy. In line with this, efficient interaction with the nuclear import machinery was shown to be a determinant of host adaptation (Gabriel et al., 2011; Hudjetz and Gabriel, 2012; Resa-Infante et al., 2008). However, the virus remains vulnerable during the transit to the nucleus, since at this stage of infection neither the classic IFN antagonist NS1 (Hale, 2014) nor the host response suppressors PA-X (Jagger et al., 2012) or PB1-F2 (Varga et al., 2011) would be expressed. Therefore, it appears plausible that the nucleocapsids themselves have to cope with the pathogen recognition system, an assumption that is in agreement with our data and also with previous reports that FLUAV polymerase subunits can prevent IFN induction (Marcus et al., 2005; Perez-Cidoncha et al., 2014) and interact with RIG-I and MAVS (Graef et al., 2010; Iwai et al., 2010; Li et al., 2014; Liedmann et al., 2014). Moreover, for influenza B virus it was demonstrated that nucleocapsids were sufficient to induce IFN, whereas for FLUAV RNA synthesis was required (Osterlund et al., 2012). These comparative data again point towards nucleocapsids being able to both induce or suppress innate immunity, dependent on the genetic background.

The connection between RIG-I activation and IFN induction is not straightforward. Avian FLUAV strains that strongly activated RIG-I were in fact weaker or equal IFN inducers than the mammalian strains whose nucleocapsids had inhibited RIG-I. Most likely, the underlying reason is found in the differences in RNA polymerase activities, which are inseparable from the RIG-I inhibition capacity. While PB2-627E nucleocapsids have low RNA synthesis rates but are strong activators of RIG-I, PB2-627K nucleocapsids have a highly active polymerase
but inhibit RIG-I. Thus, the RIG-I inhibition by PB2-627K might be overwhelmed by the higher RNA synthesis rate.

It was previously stated that IFN induction by FLUAV occurs exclusively through RNA synthesis products (Killip et al., 2014; Osterlund et al., 2012). However, we observed IRF3 activation by incoming nucleocapsids even when viral transcription was shut off by ActD treatment. This appears to contradict the results by Killip et al., who had not seen any such IRF3 activation (Killip et al., 2014). However, in that study IRF3 was assayed at 8 h post-infection, a time point at which IRF3 activity may have waned, as IRF3 returns to the inactive state in case of low inducers (Long et al., 2014). We therefore propose that incoming FLUAV nucleocapsids can activate RIG-I and antiviral signaling, similar to those of influenza B virus (Osterlund et al., 2012) and the cytoplasmic RNA viruses (Weber et al., 2013). IFN induction by nucleocapsids is however comparatively weak and the subsequent RNA synthesis becomes the dominant IFN elicitor once the nucleocapsids have reached the nucleus.

Although PB2-627K had no apparent bearing on IFN induction, for PB2-627E viruses and VLPs the establishment of infection was slowed down by RIG-I. Strikingly, while deletion of RIG-I accelerated infection by avian strains, deletion of the downstream adaptor MAVS did not. Moreover, a signaling-inactive mutant of RIG-I was as potent as wt RIG-I itself in slowing down PB2-627E viruses. It is therefore likely that the binding of RIG-I to the panhandle RNA results in a direct (although transient) antiviral effect against avian strains. The mammalian-adapted nucleocapsids are more efficient in hiding their panhandle, and it could be speculated this to be due to the stronger binding of the PB2 to the NP. The signaling-independent antiviral effect may also explain why RIG-I follows the nucleocapsids into the nucleus at the late stage of infection (Li et al., 2014).

Besides PB2-627K, there might be more factors influencing the RIG-I sensitivity of nucleocapsids. The circulating 2009 pandemic pH1N1 strains have retained the PB2-627E signature, but acquired compensating second-site mutations (Mehle and Doudna, 2009;
Yamada et al., 2010). Nonetheless, our direct comparisons between variants of four FLUAV strains that differed only in position PB2-627 strongly indicate that this major adaptation is in fact a RIG-I escape mechanism. It will be interesting to see whether other adaptive mutations in the polymerase or the NP also affect RIG-I.

FLUAV epidemics feed from avian reservoirs. Wild bird species, e.g. ducks, express functional RIG-I, but domestic chicken do not (Barber et al., 2010; Kowalinski et al., 2011). Many avian FLUAV strains cause asymptomatic infection e.g. in ducks, but an acute systemic disease in chicken (Kim et al., 2009). Populations of wild birds are comparatively disperse, i.e. overt disease would reduce the chance of viral spread. Chicken, by contrast, are held under crowded conditions and with a constant, non-natural replenishment of susceptible individuals. A certain RIG-I sensitivity may allow FLUAV to persist in the wild bird reservoir. In domestic chicken, there is no selection pressure on sparing the host, and the absence of RIG-I could enable rapid viral spread. In humans, the viral transmission mode requires replication to levels causing respiratory symptoms. Hence, it could be speculated that mutations suppressing RIG-I activation are needed for efficient and sustainable infection of humans, but not in chicken (no RIG-I) or wild birds (RIG-I enables asymptomatic infection).

While RIG-I appears to be acting on incoming nucleocapsids of avian-adapted viruses, other host cell factors are likely to contribute to the PB2-directed host restriction during later stages of infection.

Applying a polymerase-disrupting peptide massively increased RIG-I activation by nucleocapsids. This uncovers the full potential of RIG-I and implies that even for avian-adapted strains only a fraction of the nucleocapsids is actually recognized. Therefore, drugs targeting the FLUAV polymerase complex will have the secondary benefit of boosting antiviral host responses.
In sum, our data implicate that incoming FLUAV nucleocapsids are prone to a direct antiviral inhibition by RIG-I, and that the degree of RIG-I sensitivity is dependent on nucleocapsid stability.

**Experimental Procedures**

**Cells and viruses**

A549, HEK 293, DF-1, MDCK II, and *Drosophila* S2 cells were cultivated as described (Weber et al., 2013). FLUAV A/PR/8/34, the recombinant strains of A/quail/Shantou/2061/2000 (H9N2) (Baron et al., 2013), A/Hamburg/05/2009 (pH1N1), A/WSN/33 (H1N1), and A/WSN/33 with Strep-tagged PB2 (Rameix-Welti et al., 2009) were grown on MDCK II cells. A/Thai/KAN-1/04 (H5N1) was grown on chicken DF-1 cells (Manz et al., 2012). All viruses were entirely sequenced and confirmed to harbor only the intended mutations.

**Infections**

Cells were washed once with phosphate-buffered saline (PBS) and inoculated for 1 h at 4°C with virus dissolved in OptiMEM (Invitrogen) or 250 µl cell supernatant containing VLPs. The inoculum was replaced with DMEM containing 0.2% BSA and (except for VSV-G VLP infections) 1 µg/ml L-1-tosylamido-2-phenylethyl chloromethyl ketone (TPCK)-treated trypsin (Sigma-Aldrich), and cells were further incubated at 37°C.

**RIG-I activation assays**

Analyses of RIG-I conformation and oligomerization were described elsewhere (Weber et al., 2013; Weber and Weber, 2014a). For co-sedimentation assays from mammalian cells, cell
lysates were prepared as for co-immunoprecipitation (see below). The cleared lysates were loaded on a discontinuous 5% to 15% CsCl gradient in 20 mM Tris/HCl pH 7.9, 200 mM NaCl and centrifuged at 52,000 rpm for 2 h at 12°C in a SW60 rotor (Beckman). Altogether 12 fractions were recovered from top to bottom and pelleted at 45,000 rpm for 1 h at 4°C in a TLA45 rotor (Beckman). Pellets were dissolved in sample buffer, boiled for 5 min and analyzed by immunoblotting. Proteins were detected with rabbit polyclonal anti-A/quail/Shantou/2061/00 (H9N2) (Baron et al., 2013) at 1:8000, and the mouse monoclonals anti-RIG-I ALME-1 (Enzo Life Sciences; 1:1000), anti-MAVS (Abcam; 1:500), and anti-human TRIM25 (BD Transduction laboratories, 1:5000).

Co-sedimentation assays with S2 cell samples were performed by a similar procedure. Briefly, 50 µl dialyzed lysates (containing 100 µg protein) were mixed with 50 µl of dialyzed PR/8/34 nucleocapsids and supplemented with 1 mM ATP. After 1 h at 37°C the samples were analyzed using a discontinuous 10% to 30% CsCl gradient.

Inhibitor treatments

CHX, CuSO₄, ActD and IVM were purchased from Sigma Aldrich, and LMB from Biomol. Cells were pretreated with CHX (50 µg/ml, stocks dissolved in DMSO), LMB (16 nM, stocks dissolved in ethanol), ActD (1µg/ml, stocks dissolved in DMSO), or IVM (50 µM, stocks dissolved in DMSO ) for 1 h before infection. Inhibitors were also included in the virus inoculum and the incubation medium. ADP●AlF₃ treatment was performed as described (Weber et al., 2013).

Immunofluorescence microscopy

For superresolution immunofluorescence by ground state depletion (GSD) microscopy, samples were prepared as described for confocal microscopy (see supplemental information) with minor modifications. As secondary antibodies donkey anti-rabbit Alexa Fluor 555
(FLUAV) and goat anti-mouse Alexa Fluor 647 (RIG-I) were used and samples were embedded in 50 mM Tris-HCl pH 8.0 dissolved in 10% Vectashield mounting medium (VectorLabs) and 90% glycerol. Analysis was performed with the Leica SR GSD microscope.

**Purification of native viral nucleocapsids**

MDCK II cells were seeded in 9 T175 cell culture flasks at a confluency of 40% and infected with FLUAV strain PR/8/34 at an MOI of 0.01 or left uninfected (control). Supernatants were harvested at 48 h p.i. and nucleocapsids were purified as described (Weber et al., 2013).

**Co-immunoprecipitation assay**

Immunoprecipitations using mouse monoclonals anti-p21 (Santa Cruz Biotechnology) and anti-RIG-I (ALME-1) were performed as described (Weber et al., 2013). Mouse monoclonal anti-NP (HB65) (Wisskirchen et al., 2011) was used at a 1:200 dilution in RIPA buffer. Mouse monoclonals anti-p21 (1:500), anti-RIG-I ALME-1 (1:1000), and rabbit polyclonal H9N2 (1:8.000) were used for immunoblotting.

**Enzymatic treatment of nucleocapsids**

Dialyzed nucleocapsids were treated with RNase A, RNase III, or SAP and incubated with dialyzed lysates from RIG-I-expressing S2 cells (supplemented with 1 mM ATP) as described (Weber et al., 2013).

**Real-time RT-PCR**

RNA was isolated from cells using the RNeasy Mini Kit (Qiagen), and 10 ng were analyzed with the one-step QuantiTect SYBR Green RT PCR kit (Qiagen) in a StepOne Real-Time-PCR-Cycler (Applied Biosystems). Cellular RNA was quantified with specific QuantiTect primers against IFN-β (QT00203763) and β-actin (QT00996415). For detection of FLUAV
segment 7 sequences, reverse transcription was performed using the Qiagen QuantiTect® Reverse Transcription Kit with the (-) strand specific forward (5’-GGA CTG CAG CGT AGA CGC TT-3’) and (+) strand specific reverse primer (5’- CAT CCT GTT GTA TAT GAG GCC CAT-3’). PCR was performed using QuantiTect® SYBR® Green PCR. Values were normalized against γ-actin using the ΔΔCT method (Livak and Schmittgen, 2001). Up-regulation of inducible genes is depicted in relation to non-stimulated, non-infected (mock) cells.

**siRNA knockdown**

A549 cells were twice reverse-transfected with predesigned siRNAs from Qiagen (AllStar Negative Control siRNA and FlexiTube siRNAs against RIG-I (SI03649037) and MDA5 mRNA (GS23586)). For each siRNA, 25 nM were diluted in 100 µl DMEM, supplemented with 1 µl Lipofectamine RNAiMAX Reagent (Invitrogen) and incubated for 15 min at room temperature. The siRNA transfection mixes were transferred into a 24 well plate, and 1×10E5 cells dissolved in 900µl DMEM 10% FCS were seeded on top. After 48 h of incubation at 37°C with 5% CO₂, cells were harvested, and 1×10E5 cells were again reverse transfected as described above and incubated for additional 24 h at 37°C with 5% CO₂.

**Generation of knockout cell lines**

Using GeneJuice reagent (Merck), 2.5×10E4 HEK 293T cells were transfected with plasmids for either a Zinc finger nuclease targeting the *RIG-I* gene or for a pair of Transcription Activator-Like Effectors to Nucleases targeting the genes for *MDA5* or *MAVS*. Two days later, 0.5 cells per well were seeded into 6 96-well plates. After two weeks, colonies were trypsinized and seeded into two replica plates. The next day, one of the replicates was lysed (0.2 mg/ml proteinase K, 1 mM CaCl₂, 3 mM MgCl₂, 1 mM EDTA, 1% Triton-X-100, 10 mM Tris pH 7.5). Genotypes were obtained by sequencing PCR amplicons covering the target
site of interest (target sequences and primer sequences available upon request). For each
target gene, a clone was selected that had all alleles disrupted.

**VLP system**

HEK293 cells were transiently transfected with plasmids in 2 µl GeneJammer (Agilent) per 1
µg DNA, and medium was changed after 4 h. VLPs for strain A/WSN/33 were generated as
described (Neumann et al., 2000). HEK293 wt cells were transfected with plasmids for VSV-
G oder FLUAV HA (1 µg), as well as M1 (2 µg), M2 (100 ng), NEP (1 µg), PB2-627E or
PB2-627K (1 µg), PB1 (1 µg), PA (100 ng), NP (1 µg), a firefly luciferase minigenome
construct (1 µg), and a Renilla luciferase construct (pRL-SV40; 75 ng). As negative control,
PB2 was replaced by additional PB1 plasmid. Two days later supernatants were collected and
treated with 25 U/ml Benzonase (Novagen) for 3 h at 37°C. Cleared cell supernatants were
used for VLP infections of HEK293 cells (pretransfected with NP, PA, PB1 and the matching
PB2) as described above. Luciferase activities were measured 48 h post transfection (VLP-
producing cells) or infection (VLP-infected cells). Firefly luciferase activity was normalized
against Renilla activity. Relative polymerase activity is depicted as fold induction with
respect to mock control.

**Peptide inhibitor treatment**

A549 cells seeded at 80% confluency in T25 flasks were washed once with PBS and infected
with A/Thai/KAN-1/04 (MOI 1) for 1 h at 4°C. Then, cells were treated with 10 ng/ml Borna-
X-Tat or PB1_{1-15} T6Y-Tat (Wunderlich et al., 2011) dissolved in medium with 0.2% BSA. At
1 h post-treatment, cells were lysed and subjected to the RIG-I conformational switch assay.
Acknowledgements

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**Figure legends**

**Fig. 1. Activation of RIG-I signaling by incoming influenza virus nucleocapsids.**

(A and B) RIG-I activity assays. A549 cells were pre-incubated for 1 h with inhibitors, inoculated with strain A/PR/8/34 (MOI 1) or left uninfected (mock) for 1 h at 4°C, incubated 1 h at 37°C, and analyzed. (A) Oligomerization assay. Lysates of cells treated with CHX (50 µg/ml), LMB (16 nM), ActD (1 µg/ml), or IVM (50 µM) were separated by native PAGE and immunostained for RIG-I. Actin served as loading control. (B) Conformational switch. Lysates as in (A) were subjected to limited trypsin digest and analyzed by RIG-I immunoblot. The Ponceau S protein stain (representative section shown) serves as loading control for the digested samples. (C) Quantification of total FLUAV segment 7 RNA by RT-qPCR. Input represents RNA amounts harvested after the 1 h inoculation period at 4°C. (D) IRF3 activation. A549 cells were pretreated with inhibitors, inoculated with strain A/PR/8/34 (MOI 1) or left uninfected (mock) for 1 h at 4°C, and incubated for 1 h under inhibitor treatment. Lysates from cells were separated by native PAGE and analyzed by immunoblotting for phosphorylated IRF3 (P-IRF3) and actin as described (Weber et al., 2013). See also Figures S1A-S1C.

**Fig. 2. RIG-I interacts with incoming influenza virus nucleocapsids and is activated in a 5’ppp-dsRNA-dependent manner.**

(A to C) CHX / LMB-treated A549 cells were infected with A/PR/8/34 (MOI 1) for 1 h. (A) Cells analyzed for RIG-I and FLUAV by 3D GSD superresolution immunofluorescence microscopy. Scale bar, 1 µm. Insets are digitally magnified and shown below the main image (taken from one individual cell). (B) Co-immunoprecipitation. Cell lysates were subjected to immunoprecipitation (IP) using antibodies against p21 (negative control), RIG-I, or FLUAV NP, and analyzed by immunoblot. Input control: 2% of the lysate. Asterisks (*) indicate
unspecific bands. (C) Co-sedimentation assay. Cell lysates were separated by a discontinuous CsCl gradient (2% lysate as input control), and fractions analyzed by immunoblotting. (D) Conditions for activation of RIG-I by nucleocapsids in vitro. Dialyzed lysate of RIG-I-expressing S2 cells was mixed with nucleocapsids of strain A/PR/8/34 (RNPs) or a control preparation (CTRL) and supplemented with 1 mM ATP. The nucleocapsids had either been pretreated with 5 µg RNase A (A), 1 U RNase III (III), or 2 U Shrimp Alkaline Phosphatase (SAP), or left untreated (-), for 1 h at 37°C. RIG-I conformational switch was assayed after 1 h of nucleocapsid co-incubation at 37°C. See also Figures S2A-S2K.

Fig. 3. Adaptive mutations in PB2 influence the activation of RIG-I by FLUAV nucleocapsids.

(A) RIG-I activation by viruses with different PB2-627 signatures. Cells were infected with strains of A/quail/Shantou/2061/00 (H9N2), A/Thai/KAN-1/04 (H5N1), A/Hamburg/05/2009 (pH1N1), or A/WSN/33 (H1N1) containing avian-signature E or mammalian-signature K at PB2-627. Infections, CHX/LMB treatment and RIG-I conformational switch testing were performed as described for 1 and 2. (B) Quantification of virus RNAs by RT-qPCR for genomic segment 7. Input represents RNA amounts harvested after the 1-h infection period. (C) Co-sedimentation assay. Lysates of cells infected with PB2 variants of A/WSN/33 (H1N1) using our standard 1 h-protocol were separated by a CsCl gradient and analyzed by immunoblotting. (D) RIG-I-dependent IFN induction by incoming nucleocapsids. A549 cells were transfected with the indicated siRNAs or a negative control siRNA (CTRL). A549 cells siRNA-depleted of RIG-I or MDA5 were pretreated with CHX and LMB, and infected with FLUAV strains (MOI 1) for 16 h. IFN-β mRNA levels were determined by real-time RT-PCR. See also Figures S3A-S3H.

Fig. 4. RIG-I evasion by PB2-E627K.
(A) Activity of A/WSN/33-based minireplicon systems containing PB2-627K, -627E, or no PB2 (-) in HEK293 wt, ΔRIG-I or ΔMDA5 cells. (B) Reporter activities in HEK293 cells producing VLPs containing nucleocapsids with the indicated PB2 signatures. (C) Reporter activities in wt and deletion cells infected with VLPs. Cells had been pretransfected with PB1, PA, NP, and matching PB2. (D) Multicycle virus kinetics. 293 wt or ΔRIG-I cells were infected with the indicated PB2 variants of strain A/Thai/KAN-1/04 (H5N1) at an MOI 0.0001. Virus yields were determined 24 h later by plaque assay. In all cases, mean and SDs from 3 independent experiments are shown. (E) Single-cycle virus kinetics. Cells were infected at an MOI of 1 and monitored for NP expression over time. See also Figures S4A-S4D.

**Fig. 5. Signaling-independent RIG-I effect on early infection**

Single-cycle infection kinetics of A/WSN/33 on human cells lacking MAVS (A) or RIG-I (B), or on chicken DF-1 cells that naturally lack RIG-I (C). The ΔRIG-I and the DF-1 cells were transiently transfected with plasmids encoding GFP (negative control), FLAG-RIG-I wt, or FLAG-RIG-I K270A, as indicated. Overexpression was controlled using antibodies against GFP or the N-terminal Flag tag of the RIG-I constructs. See also Figure S5.

**Fig. 6. Effect of the PB2 627 signature on protein-protein interactions.**

(A) NP immunoprecipitation. Cells were CHX / LMB treated and infected with A/WSN/33 strains carrying PB2-627K or 627E as described for 1A. Lysates were immunoprecipitated 1 h later with anti-NP and immunoblotted as indicated. Normalized quantifications of the immunoprecipitated proteins are shown below. Note that amounts of viral input proteins are too low to be detected in the total extracts. (B) RIG-I immunoprecipitations from infected cells. HEK293 cells were infected with A/WSN/33 PB2 variants as described for 1A. Immunoprecipitations with anti-RIG-I and immunoblotting were performed as indicated for
2B. (C) RIG-I immunoprecipitations of recombinant nucleocapsids. HEK293 cells were transfected with A/WSN/33 NP combined with GFP or the PB2 variants (left panel), or with all A/WSN/33 minireplicon plasmids (right panel). Immunoprecipitations with anti-RIG-I were performed as indicated for (B). (D) Polymerase destabilization. A549 cells were infected with PB2 variants of strain A/WSN/33 (MOI 1), treated with peptides Borna-X-Tat (CTRL) or PB1-15 T6Y-Tat (PB1-T6Y), and tested for RIG-I conformational switch 1 h post-infection. See also Figures S6A-S6C.
Figure 2
Figure 3

Click here to download high resolution image
Figure 4

(A) Minireplicon system

PB2 plasmid:
- - 627K 627E

HEK 293: wt ΔRIG-I ΔMDA5

(B) VLP-producing cells

VLP plasmids incl.
- - PB2 627K PB2 627E

(C) VLP-infected cells

VLPS:
- - PB2 627K PB2 627E

HEK293: wt ΔRIG-I ΔMDA5 ΔMAVS

(D) Virus yields

Viruses:
- - PB2 627K PB2 627E

HEK293: wt ΔRIG-I

(E) H5N1

PB2-627E

mock 2 h 4 h 6 h 8 h

PB2-627K

2 h 4 h 6 h 8 h

time p.i.

NP

actin

HEK293

ΔRIG-I

NP

actin
Figure S1. Immediate-early activation of RIG-I is independent of viral RNA synthesis (related to Figure 1).

(A) Effect of inhibitors on viral RNA synthesis. Cells pretreated with the inhibitors used for the experiment in figure 1 were infected with FLUAV strain A/PR/8 for 8 h. The amounts of viral (-) sense RNA (segment 7) were then measured by RT-qPCR. (B) Normalization of the RNA values shown in Fig.1A. The graph illustrates the absence of RNA synthesis during the 1 h-infection period. (C) Inhibitors of translation and viral RNA synthesis do not influence immediate early activation of RIG-I. Conformational switch assay of infected A549 cells pretreated with CHX (50 µg/ml) in combination with either 16 nM LMB, 10 µM Brequinar (BRQ; depletes pyrimidine synthesis), 10 µM mycophenolic acid (MA; depletes GTP), 10 µM Pyrazofurin (PYF; depletes CTP and UTP), or 5 µM cyclopentenylcytosine (CPEC; depletes CTP).
(A) Colocalization analysis. A549 cells were pretreated for 1 h with CHX and LMB, inoculated with strain A/PR/8/34 (MOI 1) for 1 h at 4°C, and incubated for another hour at 37°C under inhibitor treatment. Cells were fixed and stained for double immunofluorescence analysis using antisera against RIG-I (red channel) and FLUAV NP (green channel). Cell nuclei were counterstained with DAPI (blue channel). The marked square area is shown digitally magnified in the lower left corner. Four fluorescence intensity overlays are shown on right. (B) Interaction of RIG-I with FLUAV nucleocapsids is independent of viral RNA synthesis. Co-immunoprecipitation analysis as performed for figure 2B, but with 1 µg/ml ActD added to the inhibitor mix containing CHX and LMB. Lysates of inhibitor-pretreated and 1 h-infected A549 cells were subjected to immunoprecipitation (IP) using antibodies against cyclin-dependent kinase inhibitor p21 (negative control), RIG-I, or FLUAV NP, and analyzed by immunoblot. Input control: 2% of the lysate. Asterisks (*) indicate unspecific bands. (C, D, E) Activation of RIG-I by nucleocapsids in the absence of RIG-I ATPase activity. Lysates from cells infected as indicated for (A) were treated with ADP●AlF3 and tested for conformational switching (C), oligomerization (D), and fractionation in a discontinuous CsCl gradient (2% lysate as input control) (E). Analyses were performed as described for figures 1 and 2. (F, G, H) FLUAV nucleocapsids directly activate RIG-I in vitro. Cell lysates of RIG-I-expressing Drosophila S2 cells were dialyzed, mixed with purified nucleocapsids (A/PR/8 RNPs) or a control preparation (CTRL), supplemented with 1 mM ATP, and assayed by limited trypsin digestion (F), native PAGE (G), or CsCl fractionation (H), coupled to immunoblot analysis. (I, J, K) RIG-I activation does not depend on the nucleocapsid preparation method. (I) Nucleocapsids of strain A/PR/8/34 were prepared and purified using a CsCl gradient (left panel) or a glycerol gradient (right panel), and their composition monitored by PB2 and NP immunoblotting (upper panels) and by Coomassie staining (lower panel). The fractions with the subsequently used nucleocapsids (RNPs) are indicated. (J) Nucleocapsids prepared using either CsCl or glycerol gradients were tested for their ability to activate RIG-I conformational switching in the insect cell / in vitro reconstitution system as described for (F). (K) Nucleocapsids of strain A/WSN/33 carrying a C-terminal strep-tag on the PB2 protein were (Rameix-Welti et al., 2009) were prepared by affinity purification and used to activate RIG-I conformational switching in the insect cell / in vitro reconstitution system. The nucleocapsids had either been pretreated with RNase A (A), RNase III (III), or Shrimp Alkaline Phosphatase (SAP), or left untreated (-) as described for Fig. 2D.
Figure S3

A

B

Ratio RNA (1 h p.i.) RNA (input)

C

D

Ratio RNA (1 h p.i.) RNA (input)

E

F

VLP envelope protein

G

H

VSV-G

WSN-HA

CTRL WSN (H1N1) WSN

mock A/PR/8 WSN (H1N1)

PB2 627

RIG-I

30 kDa RIG-I fragment Ponceau S

trypsin +

PB2 627

CTRL WSN (H1N1) WSN

mock CHX/LMB CHX/LMB + ActD

PB2 627

RIG-I

30 kDa RIG-I fragment Ponceau S

trypsin +

CTRL MDA5 Ld RIG-I Ld

mock
Figure S3. RIG-I activation by PB2-627E signature virus is mediated by nucleocapsids and independent of viral RNA synthesis (related to Figure 3).

(A) Quantification of RIG-I conformational switch provoked by different FLUAV strains. Cells were treated and infected as described for figure 3A. Signals for the trypsin resistant RIG-I fragment were quantified and normalized to the mock control. Data of the experiment shown in Figure 3A and of two other independent experiments were used to calculate values of mean and SD. (B) Normalization of the RNA values shown in Figure 3B. The quantification illustrates that nucleocapsids of avian and human strains contain similar amounts of RNA and that no RNA synthesis occurs during the 1 h infection period. (C and D) Systematic comparison of RNA amounts from input and 1 h-infections. Mean values and standard deviations from 3 independent experiments are shown (C) and normalized (D). P values were determined by Student’s t-test. (E) RIG-I activation by nucleocapsids of strain A/PR8/34 is in the range expected for a virus with a PB2-627K signature. Cells pretreated with CHX and LMB were infected for 1 h with the indicated viruses and tested for RIG-I conformational switching. (F) RIG-I activation is independent of the viral envelope. Cells pretreated with CHX and LMB were infected for 1 h with VLPs derived from strain A/WSN/33 (H1N1) that either contain the viral HA protein or that are pseudotyped with VSV-G, and tested for RIG-I conformational switching. (G) RIG-I activation by PB2-627 variant viruses is independent of viral RNA synthesis. RIG-I conformational switch assay for cells infected with A/WSN/33 (H1N1) which had been pretreated with CHX and LMB (left panel), or in addition with ActD (right panel). (H) Control of siRNA treatment efficiency. A549 cells were transfected with the indicated siRNAs or a negative control siRNA (CTRL). After two rounds of transfections, cells were tested for the presence of the target proteins by immunoblot.
Figure S4

A. Western blot analysis showing RIG-I, MDA5, MAVS, and actin levels in control (CTRL), RIG-I (ΔRIG-I), MDA5 (ΔMDA5), and MAVS (ΔMAVS) samples. The time points are indicated as 2 h, 4 h, 6 h, and 8 h p.i.

B. Western blot analysis of PB2 627E and 627K VLPs in control (CTRL), K, and E samples. The Western blot shows PB2 and actin levels.

C. Western blot analysis of H9N2 virus-infected HEK293 cells showing NP and actin levels. The time points are indicated as 2 h, 4 h, 6 h, and 8 h p.i.

D. Western blot analysis of WSN (H1N1) virus-infected HEK293 cells showing NP and actin levels. The time points are indicated as 2 h, 4 h, 6 h, and 8 h p.i.

E. A bar graph showing the log10 fold change in NP levels in WSN PB2 627K and WSN PB2 627E. The P-values are indicated as 0.97, 0.96, 0.96, and 0.87 for input and 1 h p.i. samples.
Figure S4. Onset of FLUAV infection in dependency of RIG-I (related to Figure 4).

(A) Characterization of knockout cells. Immunoblot analysis of HEK293 cells which had been engineered to disrupt the coding sequences of RIG-I, MDA5, and MAVS respectively (see Materials and Methods).

(B) PB2 of VLP-producing cells. Immunoblot analysis of HEK293 wt cells transfected to produce VSV-G pseudotyped VLPs.

(C, D) Onset of infection kinetics. HEK293 wt or ΔRIG-I cells were infected with PB2 variants of strain A/quail/Shantou/2061/2000 (H9N2) (C) or A/WSN/33 (H1N1) (D) at an MOI of 1, and tested by immunoblotting for the presence of NP at the indicated time points.

(E) Systematic comparison of RNA amounts from input and 1 h-infections of different PB2 signature viruses. Mean values and standard deviations from 3 independent experiments are shown. P values were determined by Student’s t-test.
Figure S5

Figure S5. Signaling-independent RIG-I effect on infection with PB2-627E VLPs (related to Figure 5)

Activities of PB2 variant VLPs of strain A/WSN/33 (VSV-G pseudotyped) in ΔRIG-I cells transcomplemented with GFP, wt RIG-I, or RIG-I K270A as described for Figure 5. Cells had been additionally pretransfected with PB1, PA, NP, and matching PB2. Mean and SDs from 3 independent experiments are shown.
Figure S6

A

HEK293: wt \( \Delta \text{RIG-I} \) \( \Delta \text{MDA5} \)

WSN NP, PA, PB1, vRNA-FF

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HEK293: wt \( \Delta \text{RIG-I} \) \( \Delta \text{MDA5} \)

H9N2 NP, PA, PB1, vRNA-FF

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Figure S6. Protein-protein interactions in dependency of RIG-I (related to Figure 6). (A and B) NP-PB2 co-immunoprecipitation assay for strains WSN (H1N1) and H9N2. WT, ΔRIG-I or ΔMDA5 cells were transfected with plasmids for NP, PA, PB1, and a firefly luciferase minigenome (vRNA-FF) in combination with either GFP or PB2-627K or PB2-627E. The viral expression plasmids were either derived from strain A/WSN/33 (H1N1) (A) or strain A/quail/Shantou/2061/00 (H9N2) (B). At 24 h post-transfection, cell lysates were immunoprecipitated with a mouse monoclonal antibody recognizing NP and tested by Western blotting for the presence of NP and PB2. (C) Influence of polymerase destabilization on RIG-I activation by strain H5N1. A549 cells were pretreated with 10 ng/ml of peptides Borna-X-Tat (CTRL) or PB1_{1-15} T6Y-Tat (PB1-T6Y) and infected with A/Thai/KAN-1/04 (H5N1) strains of the specified PB2 types. At 1 h post-infection, cells were lysed and subjected to limited trypsin digestion.
Supplemental Experimental Procedures

Inhibitor treatments

Brequinar (BRQ) and mycophenolic acid (MA) were purchased from Sigma Aldrich. Pyrazofurin (PYF, NSC-143095) and cyclopentenylcytosine (CPEC, NSC-375575) were kindly obtained from the Drug Synthesis and Chemistry Branch of the National Cancer Institute.

Cells were pretreated for 24 h before infection with CHX (see main text) combined with BRQ (10 µM, stocks dissolved in DMSO), MA (10 µM, stocks dissolved in methanol), PYF (10 µM, stocks dissolved in DMSO) or CPEC (5 µM, stocks dissolved in DMSO). Inhibitors were also included in the virus inoculum and the incubation medium.

Preparation and transfection of virus particle RNA

MDCK II cells grown in a T175 flask were infected with A/PR/8/34 (MOI 0,001). At 38 h post infection, supernatants were collected and centrifuged at 1,500 rpm for 7 min at 4°C in a JA-12 rotor (Beckman). The cleared supernatant was supplemented with 5 M NaCl and 30% PEG800 in NTE buffer (10 mM Tris-HCl pH 6.5, 1 mM EDTA, 100 mM NaCl) to a final concentration of 0.44 M NaCl and 15% PEG800 in NTE buffer, respectively. The suspension was incubated on ice for 30 min under constant shaking and centrifuged at 6,000 rpm for 1 h at 4°C. Subsequently, the pellet was resuspended in 1.5 ml peqGOLDTriFAST (Peqlab). After 5 min incubation at room temperature, 300 µl chloroform were added and the suspension was incubated for another 10 min at room temperature. After centrifugation at 4,600 rpm for 11 min at 4°C, the aqueous phase was collected and supplemented with 30 µg glycogen (Roche Diagnostics) and one volume of isopropanol. The RNA was precipitated for 16 h at -20°C and then pelleted at 4,600 rpm for 30 min at 4°C. The pellet was washed twice by adding 2 ml 70% ethanol followed by centrifugation at 12,000 × g for 10 min at 4°C. After the remaining ethanol had evaporated completely, the pellet was resuspended in TE buffer (10 mM Tris-HCl pH 8.0, 1 mM EDTA) and stored at -80°C.

For transfection, 500 ng virus particle RNA was incubated for 10 min at room temperature with 2 µl/µg RNA jetPRIME (PolyPlus Transfection) according to the manufacturer’s instructions. The transfection mix was added dropwise onto human A549 cells grown in a T25 flask to 80% confluence and the cells incubated for 1 h at 37°C before lysis.
**Immunofluorescence microscopy**

For immunfluorescence analysis, human A549 cells were grown on coverslips and infected for 1 h as outlined above. Then, cells were fixed with 4% paraformaldehyde dissolved in PBS, permeabilized with 0.5% Triton X-100 dissolved in PBS for 15 min, and blocked with 2% bovine serum albumine (BSA), 5% glycerol, 0.2% Tween 20 dissolved in PBS for 1 h at room temperature. Primary antibodies, rabbit polyclonal recognizing A/quail/Shantou/2061/00 (H9N2) and mouse polyclonal anti-human RIG-I (1:200; Baum et al., 2010), were diluted in blocking solution and cells were stained for 1 h at room temperature. After three times washing with PBS, cells were incubated with Alexa Fluor 555 donkey anti-mouse (1:200) and Alexa Fluor 488 donkey anti-rabbit (1:200) secondary antibodies supplemented with 4,4-diamidino-2-phenylindole (DAPI, 1:10.000) for 45 min at room temperature. Cells were washed three times with PBS and coverslips were mounted using Fluorsave solution (Calbiochem). Stained cell samples were examined using a Leica SP5 confocal microscope.

**Expression of RIG-I in S2 cells**

*D. mel.* S2 cells were transfected with 1 µg of pRmHa3-RIG-I plasmid, induced with CuSO₄, lysed and extracts dialyzed as described (Weber et al., 2013).

**Purification of recombinant viral nucleocapsids**

For purification of recombinant nucleocapsids via Strep-affinity tag, MDCK II cells in a 10 cm dish were infected with A/WSN/33 carrying a C-terminal Strep-tag on the PB2 protein (Rameix-Welti et al., 2009) at an MOI of 1. After 16 h the cells were washed with PBS, scraped off in PBS, pelleted by centrifugation at 100 × g for 5 min at 4°C. Pellets were then resuspended in Strep-lysis buffer (50 mM Tris/HCl pH 7.5, 150 mM NaCl, 2 mM MgCl₂, 2mM CaCl₂, 1% PMSF, 1 mM DTT, 0.7% NP40 and 50 U/ml DNase I (Fermentas)) and incubated on ice for 20 min. Lysates were then centrifuged at 16.200 x g for 10 min at 4°C and the cleared lysates were incubated for 1 h at 4°C with Strep-Tactin beads (IBA Goettingen). Beads were then pelleted by centrifugation at 100 × g for 1 min at 4°C, supernatant was removed and the beads were resuspended with wash buffer (50 mM Tris/HCl pH 7.5, 150 mM NaCl, 2 mM MgCl₂, 2mM CaCl₂, 1% PMSF, 1 mM DTT). Washing was performed three times and nucleocapsids were eluted by incubating the beads for 15 min on ice with wash buffer supplemented with 2.5 mM desthiobiotin (IBA Goettingen) Suspension was centrifuged at 100 × g for 1 min at 4°C and the eluate was transferred in a fresh tube and nucleocapsids were dialyzed against PBS.
Co-immunoprecipitation assay of recombinant nucleocapsids

For immunoprecipitations of recombinant nucleocapsids produced in minireplicon systems of strains A/WSN/33 (Neumann et al., 2000) or H9N2 (Baron et al., 2013), cells were seeded in T25 flasks and transiently transfected with expression plasmids encoding GFP (1 µg) or strain-specific PB2-627K or 627E (1 µg) together with NP (1 µg) alone or additionally with strain-specific PB1 (1 µg), PA (100 ng) and the firefly luciferase reporter minigenome construct (1 µg). At 24 h post-transfection, cell lysates were prepared and subjected to co-immunoprecipitation assay. Thereby, beads were directly coupled with mouse monoclonal HB65 antibody recognizing NP (Wisskirchen et al., 2011) or RIG-I (ALME-1). Coupled beads were incubated with the lysates for 2 h at 4°C and eluted as described. Co-immunoprecipitations were validated by immunoblotting using rabbit polyclonal H9N2, mouse monoclonal anti-PB2 JG6 (Marazzi et al., 2012) and mouse monoclonal anti-RIG-I antibody ALME-1.
Supplemental Reference

4.4. Intergenic region of incoming nucleocapsid activate PKR
-Lassa virus nucleoprotein provides a PKR evasion strategy-

Own contribution:

I performed all experiments of the main figures. Samples of Suppl. Fig. 2 were prepared by Markus Kainulainen (group of Prof. Dr. Weber, Institute for Virology, Marburg), Sarah Katharina Fehling (group of Dr. Strecker, Institute for Virology, Marburg) and Svenja Wolff (group of Prof. Dr. Becker, Institute for Virology, Marburg) while I performed analysis of the samples. I contributed to writing of the manuscript.

Michaela Gerlach
Intergenic region of incoming nucleocapsid activate PKR -Lassa virus nucleoprotein provides a PKR evasion strategy-

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Abstract

Surveillance of non-self RNA by receptors of the innate immune system is a crucial first line of defense against viral infection. A rapid detection is thereby required for initiating an antiviral response to ensure survival of the host. Protein kinase R (PKR) represent one immune sensor able to recognize dsRNA and specific RNA secondary structures. Despite recognizing viral replicative dsRNA intermediates, we identified in this study the hairpin-structured intergenic region (IGR) of viral ambisense genomes as natural PKR agonists. Thus, the IGR seems to be exposed from nucleoprotein protection within the nucleocapsid complex allowing PKR to interact. Upon IGR recognition, PKR undergoes conformational switching and phosphorylation, identifying IGRs of RVFV S segment, New World arenavirus Tacaribe (TCRV) and Junin virus (JUNV) and Old World arenavirus Lassa virus (LASV) nucleocapsids as natural PKR agonists. To antagonize antiviral activity of PKR, TCRV, JUNV and LASV induce proteasomal degradation of activated PKR. In case of LASV, the newly synthesized nucleoprotein interacts with PKR and promotes its degradation. Hence, we identified the IGR as part of the viral nucleocapsid as an immediate PKR agonist. Furthermore, we pin down LASV nucleoprotein as a PKR antagonist by promoting its proteasomal degradation. This highlights PKR as an immune sensor of immediate infection and that arenaviruses have adapted to rapidly and efficiently avert antiviral PKR activity. Furthermore, IGRs play an essential role in the control of viral protein synthesis indicating the necessity to conserve this structure despite the presence of antiviral defense mechanisms. This suggests targeting of the IGR to block virus replication a promising target for drug development.
Introduction

Virus infection is a constant threat to human mankind. Thus, to rapidly control virus infection an immediate recognition of the viral intruder is required. Several groups of innate immune receptors, including Toll-like receptors (TLR), RIG-I like receptors (RLR), NOD-like receptors (NLR), C-type lectin receptors (CLR), and other cytosolic RNA and DNA sensors are thereby crucial to distinguish between host cell (self) and pathogen associated structures (non-self) (Pandey et al., 2014).

RLR have a central role to detect RNA virus infection in the cytoplasm of the infected cell. The family of RLR is formed by retinoic acid inducible gene 1 (RIG-I), melanoma differentiation association factor 5 (MDA5) and laboratory of genetics and physiology 2 (LGP2). Studies have demonstrated that RIG-I and MDA5 recognize mostly distinct RNA structures. MDA5 is known to sense longer double-stranded RNA (dsRNA) molecules, ideally with higher order RNA structures (Pichlmair et al., 2009, Schlee, 2013). RIG-I mainly responds to 5’ ppp blunt-ended dsRNA of a minimal length of 10 base pairs (Kohlway et al., 2013, Schmidt et al., 2009, Schlee et al., 2009). Recently, dsRNA stretches bearing a 5′diphosphate were identified to stimulate a RIG-I dependent response (Goubau et al., 2014). Interestingly, also in the encapsidated state, RIG-I can sense short 5′ppp dsRNA panhandle structures of incoming nucleocapsids (Weber et al., 2013). Ligand-mediated activation of RIG-I stimulates a signaling reaction that culminates in the induction of type I interferon (IFN-alpha/beta), proinflammatory cytokines and eventually numerous interferon stimulated genes (ISGs) (Goubau et al., 2013, Reikine et al., 2014, Ivashkiv and Donlin, 2014, Hertzog and Williams, 2013). ISG represent viral restriction factors that directly inhibit all steps of virus replication and furthermore co-stimulatory molecules, cytokines and chemokines which favor the initiation of adaptive immune responses (Schoggins, 2014, Schneider et al., 2014, Loo and Gale, 2011).

Viruses have evolved to impair or evade RIG-I recognition. Further immune sensors are thus required to ensure a rapid detection and response to the viral intruder. However, whether other immune sensors recognize like RIG-I incoming protein-associated viral RNA and stimulate an antiviral response remained elusive. Here, we addressed the question whether protein kinase R (PKR) contributes to an immediate immune recognition.

PKR represents an RNA sensor which is expressed at a low constitutive level mainly in the cytoplasm. Despite being upregulated upon virus induced IFN response, PKR expression can also be increased upon other cellular stress signals like disturbances of calcium homeostasis (Williams, 1999). PKR represents a multifunctional protein involved in the regulation of cap-dependent translation by phosphorylation of the alpha subunit of eukaryotic initiation factor 2 (eIF2α). Furthermore, PKR is crucial for the virus induced formation of stress granules required for IFN-signaling (Onomoto et al., 2012). To stimulate PKR activation there is a minimal dsRNA length requirement of approximately 33 base pairs, but also specific structural modalities like bulges and loops and nucleotide modifications
like a 5′ppp support PKR stimulation (Nallagatla et al., 2007, Dzananovic et al., 2013, Bevilacqua and Cech, 1996). Thereby, replicative intermediates of negative-strand RNA viruses can form PKR activating dsRNA stretches. PKR thus senses virus replication, but involvement of PKR as an immediate immune sensor and inducer of antiviral defense mechanisms remains to be resolved. An interaction of PKR with viral RNA secondary structures like the hepatitis C virus IRES or dimerized stem-loop structures of transactivation (TAR) RNA of human immunodeficiency virus 1 was previously reported (Shimoike et al., 2009, Heinicke et al., 2009). Whether PKR would however recognize viral RNA in the encapsidated state was not addressed so far.

In this study, we report the hairpin-structured intergenic region (IGR) of two distinct virus families as PKR agonists. Thereby, the IGR seems to be exposed from nucleoprotein protection within the nucleocapsid complex allowing PKR to interact. Upon IGR recognition, PKR undergoes conformational switching and phosphorylation identifying IGRs of RVFV S segment, New World arenavirus Tacaribe (TCRV) and Junin virus (JUNV) and Old World arenavirus Lassa virus (LASV) nucleocapsids as natural PKR activators. To antagonize antiviral activity of PKR, TCRV, JUNV and LASV induce proteasomal degradation of activated PKR. In case of LASV, the newly synthesized nucleoprotein interacts with PKR and promotes its degradation. Hence, we identified the IGR as part of the viral nucleocapsid as an immediate PKR agonist. LASV nucleoprotein counteracts antiviral PKR activity by promoting its proteasomal degradation. Furthermore, this seems to be a common mechanism within the arenavirus family although the responsible TCRV and JUNV protein could not be identified so far. This highlights PKR as an immune sensor of immediate infection and that arenaviruses have adapted to rapidly and efficiently avert antiviral PKR activity.
Experimental Procedures

Cells, Viruses and Transfection

Human A549 and HEK293T cells were cultivated in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal calf serum (FCS) at 37°C and 5% CO₂. Recombinant La Crosse virus lacking the virulence factor NSs (LACV; (Blakqori and Weber, 2005)) and recombinant Rift Valley fever virus expressing Renilla luciferase instead of NSs (RVFV; (Habjan et al., 2009a)), Junin virus (JUNV), Tacaribe virus (TCRV) and Lassa virus (LASV) were propagated on Vero E6 cells. All work with LASV and JUNV was performed in the BSL-4 laboratories at the Philipps University Marburg, Germany.

For transient overexpression, HEK293T cells were transfected by standard calcium phosphate transfection method with either 0.25, 0.5, 1 or 2 µg of pCAGGS_GP-HA, pCAGGS_NP-HA, pCAGGS_Z-HA or pcDNA3.1_ΔMX combined with 1 µg of pcDNA5_PKR.

Rift Valley Fever virus like particle (RVF VLP) production

Generation of RVF VLPs was carried out as previously described (Habjan et al., 2009a), with minor alterations. Briefly, 1x10⁶ HEK293T cells were transfected with 1 µg each of plasmid pl.18_RVFV_N, pl.18_RVFV_L, pl.18_RVFV_M, and a reporter minigenome using Nanofectin transfection reagent (PAA Laboratories). Therefore, a minigenome with Renilla luciferase in negative sense flanked by 5’ and 3’ untranslated regions (UTRs) from either RVFV middle (M) or small (S) segment was used (pHH21_RVFV_vM and pHH21_RVFV_vS, respectively). In case of the S segment, the minigenome contained the full-length RVFV S segment and the NSs replaced by Renilla luciferase. For production of empty VLPs (no minigenome), 1 µg each of plasmid pl.18_RVFV_N, pl.18_RVFV_M and for DNA amount normalization pcDNA3.1_ΔMX was used for transfection. At 4 h post-transfection, the medium was exchanged and 48 h later supernatants were collected and clarified from cell debris by centrifugation at 500xg for 5 min. Subsequently, VLPs were treated with 25 U/ml benzonase (Novagen) for 3 h at 37°C and then stored at -80°C.

Infection and Inhibitor treatment

A549 cells were inoculated for 1 h with viruses or RVF VLPs. Infection medium was replaced with DMEM containing 2% FCS thereafter. If indicated, cells were pretreated with 50 µg/ml cycloheximide (CHX) dissolved in complete medium for 1 h. CHX was also added to the virus inoculum and to the medium added afterwards. MG132 dissolved in media at the final concentration of 20 µM was added immediately after the virus inoculum was removed. In case of transfected cells, 20 µM MG132 containing complete media was added 8 h post-transfection and cells were incubated for additional 16 h at 37°C with 5% CO₂. CHX and MG132 were purchased from Sigma-Aldrich. Cells were lysed or fixed at the indicated time points.
Limited protease digestion

Limited protease digestion to monitor RIG-I and PKR conformational switching was essentially performed as described elsewhere (Weber and Weber, 2014a). Briefly, cells of one well of a six well plate were lysed in 25 µl 0.5% Triton X-100 in PBS supplemented with Phosphatase Inhibitor Cocktail set II (Calbiochem). After an incubation on ice for 10 min, the samples were centrifuged for 10 min at 10,000×g at 4°C and 10 µl of the cleared lysate were kept as input and the other 10 µl were digested for 20 min with 0.2 µg/µl L-1-tosylamido-2-phenylethyl chloromethyl ketone (TPCK)-treated trypsin (Sigma-Aldrich) at 37°C. Reaction was stopped by adding 4-fold sample buffer (200 mM Tris-HCl pH 6.8, 8% SDS, 40% glycerol, 25% β-Mercaptoethanol, 0.4% Bromphenol Blue) and boiling for 10 min at 100°C according to BSL4 safety procedures. Samples were subjected to SDS-PAGE and Western Blot analysis using mouse monoclonal anti-RIG-I antibody (ALME-1, Enzo Life Sciences) at 1:1,000; mouse monoclonal anti-PKR B10 (Santa Cruz) at 1:500 and rabbit monoclonal against phosphorylated PKR at position threonine 446 (p-PKR Thr 446; Epitomics) at 1:500.

Co-immunoprecipitation

Cells grown in a T75 tissue culture flask (infection) or in two wells of a six well plate (transfection) were scraped off in PBS, centrifuged at low speed, and the pellets were lysed in 50 mM Tris-HCl pH 7.5, 150 mM NaCl and 1% NP40 supplemented with phosphatase inhibitor and Protease inhibitor cocktail Complete (Roche), incubated on ice for 10 min and centrifuged for 10 min at 10,000×g at 4°C. Cleared cell lysates were transferred to fresh tubes and 2% were kept as input control. The remaining lysate was subjected to immunoprecipitation (IP) using magnetic beads (Dynabeads, Invitrogen). The coupling of the beads with the corresponding antibody was performed according to the manufacturer’s instructions with minor modifications. For each IP, 1.5 mg of beads were incubated with rabbit polyclonal anti-mouse antibody (DAKO) at a dilution of 1:200 for 16 h at 37°C. After the first antibody coupling, beads were washed three times with lysis buffer, and the anti-mouse antibody beads were further incubated with mouse monoclonal anti-p21 (Santa Cruz Biotechnology), mouse monoclonal anti-RIG-I or mouse monoclonal anti-PKR B10 at a 1:200 dilution in lysis buffer for 24 h at 4°C. Beads were washed three times with lysis buffer and then incubated with the lysates for 2 h at 4°C. The immunoprecipitates were washed three times with lysis buffer and eluted by adding 1-fold sample buffer (50 mM Tris-HCl pH 6.8, 2% SDS, 10% glycerol, 6.25% β-Mercaptoethanol, 0.1% Bromphenol Blue). Western blot analysis was performed with mouse monoclonal anti-RIG-I at 1:1000, mouse monoclonal anti-PKR B10 at 1:500, rabbit monoclonal anti-p-PKR Thr 446 at 1:500, mouse monoclonal anti-p21 (Santa Cruz) at 1:500, rabbit polyclonal anti-HA (abcam) at 1:1000, rabbit polyclonal anti-LACV N at 1:1000, mouse polyclonal anti-RVFV N serum (Habjan et al., 2009b) at 1:1000 and rabbit monoclonal anti LASV NP at 1:3000. As a secondary antibody protein A-horseradish
peroxidase (HRP) conjugate (Millipore) at a dilution of 1:10,000 was used to avoid detection of heavy and light antibody chains.

**Co-sedimentation assay**

For co-sedimentation assays, cell lysates were prepared as described for co-IP. The cleared lysate was loaded on top of a discontinuous 50% to 70% glycerol gradient in 50 mM Tris-HCl pH 7.5, 150 mM NaCl and centrifuged at 52.000 rpm for 2 h at 12°C in a SW60 rotor (Beckman). Twelve fractions were recovered from top to bottom and pelleted at 45,000 rpm for 1 h at 4°C in a TLA45 rotor (Beckman). Pellets were dissolved in 1-fold sample buffer, boiled for 5 min at 100°C and validated by SDS-PAGE followed by Western Blot analysis. Proteins were detected with mouse monoclonal anti-RIG-I antibody at 1:1000, mouse monoclonal anti-PKR B10 at 1:500, rabbit polyclonal anti-LACV N at 1:1000, and mouse polyclonal anti-RVFV N serum at 1:1000.

**GSDIM**

For immunofluorescence analysis via ground state depletion microscopy followed by individual molecule return (GSDIM), 4x10e5 A549 cells were grown on coverslips and infected for 1 h as outlined above. Then, cells were fixed with 4% paraformaldehyde dissolved in PBS for 20 min, permeabilized with 0.5% Triton X-100 in PBS for 15 min, and blocked with 2% bovine serum albumine, 5% glycerol, 0.2% Tween 20 in PBS for 1 h. Primary antibodies, rabbit anti-RVFV MP12 hyperimmune serum C2, mouse polyclonal anti-human RIG-I (Baum et al., 2010) and mouse monoclonal anti-PKR B10, were diluted 1:200 in blocking solution and cells were stained for 1 h. After three times washing with PBS, cells were incubated with secondary antibody Alexa Fluor 555 donkey anti-mouse at 1:200 and Alexa Fluor 647 donkey anti-rabbit at 1:200 supplemented with 4,4-diamidino-2-phenylindole (DAPI) at 1:10,000 for 45 min at room temperature. Cells were washed three times with PBS and samples were embedded in freshly prepared 100 mM β-mercaptoethylamine in PBS pH 7.4 directly before imaging. Analysis was performed with the Leica SR GSD microscope. Quantification was performed of three independent experiments. Student’s T-test was carried out for statistical analysis.
Results

RIG-I and PKR differ in their agonist specificity

It was recently described by our group that the 5’ppp panhandle structure of incoming viral nucleocapsids serve as a natural RIG-I agonist to promote an immediate antiviral IFN response (Weber et al., 2013). To test whether other immune receptors would contribute to an immediate virus recognition, cells were infected with either La Crosse virus (LACV) or Rift valley fever virus (RVFV) (Bunyaviridae), both lacking the virulence marker NSs. Bunyavirus are peculiar since their transcription is dependent on an on-going translation (Elliott, 1990). Hence, treatment with the translation inhibitor cycloheximide (CHX) aborts viral transcription and allows analysis of immune receptor activation by incoming nucleocapsids in the physiological context of virus infection.

By performing limited protease digestion to monitor RIG-I activation status (Weber and Weber, 2014a), stimulated RIG-I undergoes conformational rearrangements leading to detection of trypsin resistant fragments. RIG-I activation is thereby independent whether viral replication occurs (Fig. 1A, left panel) or if only incoming nucleocapsids are present (Fig. 1A, right panel).

PKR is another well-known cytoplasmic pattern recognition receptor able to detect dsRNA stretches of at least 33 nucleotides or RNA secondary structures (Dabo and Meurs, 2012). PKR is composed of two amino-terminal dsRNA binding domains (dsRBD), a carboxy-terminal serine/threonine kinase domain and both domains are linked by a flexible linker (VanOudenhove et al., 2009). In absence of a stimulus, PKR is present as a monomer with an extended open conformation where the dsRBD are impeding kinase domain activity. Agonist recognition induces conformational rearrangements to a closed conformation allowing dimerization and auto-phosphorylation (Dabo and Meurs, 2012, Clemens, 1997, VanOudenhove et al., 2009, Lemaire et al., 2006). Like RIG-I, also PKR conformational changes can be measured by limited protease digestion (Weber and Weber, 2014a). Upon agonist recognition, PKR reorganizes its domains leading to partial protease resistance. Surprisingly, LACV and RVFV differ in their ability to promote PKR resistance to trypsin treatment and induction of PKR phosphorylation at threonine 446. LACV full viral replication only poorly and incoming LACV nucleocapsids even fail to induce these two markers of PKR activation (Fig. 1B). However, RVFV full infectious cycle and incoming nucleocapsids lead to a robust induction of PKR resistant fragments and detection of PKR phosphorylation. Since strongest differences between LACV and RVFV induced PKR activation were observed by stimulating with incoming nucleocapsids, we focused on the immediate pathogen recognition.

By performing co-sedimentation assay the migration pattern in a discontinuous glycerol gradient of RIG-I and PKR upon stimulation with LACV or RVFV nucleocapsids was analyzed. In mock infected cells, RIG-I was mainly distributed from fraction 6 to 8 (Fig. 1C, upper panel). LACV and RVFV nucleocapsids
induced a shift of RIG-I up to fraction 4 confirming the previous observation that RIG-I is able to recognize these two bunyaviruses (Fig. 1C, middle and lower panel). PKR distribution upon LACV nucleocapsid stimulation showed a comparable distribution pattern from fraction 7 to 10 as mock infection (Fig. 1C, upper and middle panel). In contrast, RVFV nucleocapsids induced a concentration of total PKR in fraction 7 (Fig. 1C, lower panel). The observed shift of nucleocapsids can hence be associated to recognition of LACV and RVFV nucleocapsids by RIG-I or RIG-I and PKR, respectively. Noteworthy, migration of RIG-I and PKR was dependent on RNA since overexpression of the nucleoprotein alone did not induce a change in the distribution pattern (Suppl. 1).

To further validate an interaction between PKR and incoming bunyavirus nucleocapsids co-immunoprecipitation (co-IP) assays were performed. As reported before, RIG-I interacted with LACV as well as RVFV nucleocapsids (Fig 1D). However, with a PKR specific co-IP only RVFV, but not LACV nucleocapsids could be co-precipitated.

In summary, these data indicate that LACV and RVFV nucleocapsids can be both recognized by RIG-I and mediate its activation. RVFV nucleocapsids, on the contrary, serve also as PKR agonist.

**Presence of an intergenic region within viral nucleocapsids determines PKR recognition and activation**

Since the observed differences of LACV and RVFV nucleocapsid recognition by PKR seemed to be dependent on viral RNA, we wondered whether this could be due to the different LACV and RVFV coding strategies. Bunyavirus particles comprise three genome segments with negative polarity which are named according to their size large (L), middle (M) and small (S). RVFV L and M segment contain only a single transcriptional unit in negativesense orientation, whereas the S segment uses ambisense coding strategy (Giorgi et al., 1991). Thereby, two open reading frames are separated by a highly conserved non-coding intergenic region (IGR). RVFV IGR forms a hairpin structure with a central dsRNA stem with internal loops and peculiar poly-C (viral sense) and poly-G (viral anti-sense) repeats (Fig. 2A) (Lara et al., 2011, Ikegami, 2012). Members of the Orthobunyavirus family like LACV do not use ambisense coding strategy (Elliott, 2014). During the replication cycle, each genome segment is transcribed into mRNA and then translated to allow genome replication and generation of progeny viruses (Fig. 2B). Amplification of the viral genome (vRNA) involves the synthesis of an exact copy of the vRNA, called complementary RNA (cRNA). In case of the RVFV S segment, the cRNA serves also as a template for mRNA synthesis (Ikegami, 2012, Walter and Barr, 2011). Since it was described that the RVFV IGR is accessible for other proteins (Moy et al., 2014) we wondered whether PKR would also use this structure to gain access to RVFV nucleocapsids.

To validate the importance of the IGR for PKR activation, virus like particles (VLPs) were generated containing nucleocapsids either with the RVFV M or S genome segment. PKR activation status was
monitored by detection of protease resistance. As a control we included RIG-I activation since 5`ppp panhandle structures of M and S segmented nucleocapsids should activate RIG-I to the same extend. Indeed, infection with RVF VLPs regardless which genome segment is packaged within the particle as well as nucleocapsids from RVFV infection stimulated RIG-I conformational rearrangements (Fig. 2C). Likewise, stimulation with RVF VLPs with S segmented nucleocapsids induce PKR resistant fragments whereas VLPs containing an encapsidated RVFV M genome segment failed to promote PKR activation. To visualize the interaction of RIG-I and PKR with the RVF VLP M or S nucleocapsids super-resolution microscopy was employed. As previously reported, RVFV nucleocapsids are displayed as pseudocircular structures with RIG-I attached via a single site (Fig 2D). Thereby, interaction of RIG-I with RVFV nucleocapsids was independent of which genome segment was packaged in the viral particle. Also PKR co-localized with nucleocapsids of RVFV infection presenting all genome segments. This interaction was significantly reduced if only M segmented nucleocapsids were present. In case of RVF VLPs enclosing S segmented nucleocapsids almost 80% of the viral nucleocapsids co-localized with PKR. Like RIG-I also PKR contacts the nucleocapsids only via a single position, most likely the IGR. This indicates that PKR recognizes the IGR of incoming RVFV nucleocapsids which promotes PKR activation.

**Arenavirus nucleocapsids promote PKR activation in absence of translation**

In comparison to RVFV, representatives of the *Arenaviridae* contain two genome segments and each uses ambisense coding strategy with an IGR separating the open reading frames (McLay et al., 2014). Arenavirus IGR represents like RVFV IGR a hairpin structure composed of a dsRNA stem with internal loops and sporadic mismatches (Fig. 3a). Therefore, we hypothesized that during arenavirus infection nucleocapsids would present strong stimulators of PKR activation. New World Junin virus (JUNV) and Tacaribe virus (TCRV) and Old World Lassa virus (LASV) were chosen as representatives of the arenavirus family to validate the potential of their nucleocapsids to promote PKR activation. Surprisingly, full infectious cycle of TCRV, JUNV and LASV did not stimulate PKR phosphorylation and conformational switching (Fig. 3b). An abort of the infectious cycle at primary transcription due to CHX treatment, however, partially rescued PKR activation. These results could be confirmed by a time-course experiment (Suppl. 2). LACV full replication cycle induced a robust PKR activation whereas LACV nucleocapsids did not stimulate a detectable amount of phosphorylated PKR. Furthermore, 24 h post-infection total PKR gets degraded indicating the induction of a negative feedback loop to control PKR signaling. In contrast, RVFV infection and incoming nucleocapsids induced phosphorylation of PKR. Therefore, only incoming nucleocapsids containing an IGR are able to promote PKR activation as observed in Fig. 1B. Full infectious cycle of TCRV, JUNV and LASV did not stimulate PKR phosphorylation though translation inhibition did partially rescue PKR activation. This indicates on the one hand, that arenavirus nucleocapsids are indeed able to stimulate PKR activation. On the other hand, PKR
activation by arenaviruses can only be detected in absence of protein synthesis. This raises the question whether newly synthesized arenavirus proteins prevents PKR activation during the full viral life cycle.

**LASV nucleoprotein induces proteasomal degradation of phosphorylated PKR**

To gain further insights into an arenavirus PKR antagonist, the focus was put on LASV. By performing co-IPs, an interaction of PKR with nucleocapsids which are just entering the cell (CHX), but also with nucleocapsids generated during viral replication (UT) could be observed (Fig. 4A). However, phosphorylated PKR could only be detected if translation was blocked. Furthermore, a weak interaction of RIG-I with LASV nucleocapsids could only be detected, when full viral replication was allowed. This is confirm with the observation that 5’ overhangs of arenavirus nucleocapsids hinder RIG-I recognition (Marq et al., 2010) and that RIG-I might rather interact with arenavirus dsRNA replicative intermediates generated during virus replication.

To identify the responsible protein able to impair PKR activation, LASV nucleoprotein (NP), glycoprotein (GP) and matrix protein (Z) were overexpressed with increasing concentrations together with PKR at a constant concentration. PKR overexpression alone was able to promote PKR phosphorylation (Fig. 4B). Thus, in case of excessive PKR within the cytoplasm of the transfected cells two PKR molecules are able to promote each other’s phosphorylation due to proximity. PKR activation status was further supported with increasing amounts of LASV GP and Z. However, overexpression of LASV NP results in a reduction of phosphorylated PKR in a dose-dependent manner, without having major effects on the total PKR amount. By performing co-IP, phosphorylated PKR is reduced in LASV NP expressing cells and also total PKR seems to be slightly decreased (Fig 4C). With a PKR specific co-IP, LASV NP, but not GP or Z, could be co-precipitated indicating an interaction between PKR and LASV NP.

As described above, RVFV and arenavirus nucleocapsids are likewise recognized by PKR. Due to this first similarity between the representatives of these two distinct virus families argues whether they would also share similar PKR antagonistic mechanisms. Hence, since RVFV NSs promotes the proteasomal degradation of PKR (Ikegami et al., 2009, Habjan et al., 2009b) we wondered whether the proteasome would also be involved in reduction of activated PKR during the course of arenavirus infection. Thus, cells were transfected with the individual LASV proteins together with PKR and analyzed for PKR phosphorylation status in absence or presence of the proteasomal inhibitor MG132. In untreated (UT) cells LASV NP overexpression reduces PKR phosphorylation in comparison to LASV GP and Z, and GFP control (Fig. 4D, left panel). However, if proteasomal degradation is blocked PKR phosphorylation is rescued in LASV NP transfected cells (Fig. 4D, right panel). However, LASV GP and Z do not affect PKR activation status in absence or presence of MG132. Also under infectious conditions,
in untreated cells (UT) TCRV, JUNV and also LASV infection failed to promote PKR activation which could be restored by MG132 treatment (Fig. 4E). This indicates that LASV NP specifically induces the proteasomal degradation of activated PKR. Additionally, since inhibition of the proteasomal pathway also rescues phosphorylated PKR in case of TCRV and JUNV suggests a similar mechanism to prevent antiviral PKR activity throughout the arenavirus family.

Taken together, after LASV entry the viral nucleocapsids are released into the cytoplasm of the infected cell for viral transcription and replication (Fig. 5). The IGR of the viral nucleocapsids is recognized by PKR, which leads to PKR conformational rearrangements and phosphorylation. Activated PKR initiates translation block via phosphorylation of eIF2α and type I IFN and proinflammatory cytokine response via NF-κB to impair LASV replication. Once inside the cytoplasm however transcription and viral protein synthesis is rapidly initiated. With an increasing concentration within the cell, LASV NP interacts with PKR and induces its proteasomal degradation. This identifies nucleocapsids bearing an IGR as natural PKR agonists and that arenaviruses have evolved to rapidly counteract immediate immune recognition.
Discussion

Surveillance of non-self RNA by receptors of the innate immune system is a crucial first line of defense against viral infection. A rapid detection is thereby required to efficiently control or even eliminate the viral intruder. Our group has recently shown, that RIG-I is able to recognize the 5′ppp dsRNA panhandle structure of incoming viral nucleocapsids to induce an immediate antiviral type I IFN response (Weber et al., 2013). However, whether other host receptors would likewise be able to recognize viral structures immediately after virus entry remained unresolved.

Thus, PKR activation by incoming nucleocapsids of two representatives of the Bunyaviridae was validated. Surprisingly, PKR recognizes and gets activated by RVFV but not LACV nucleocapsids. Considering the differences of PKR activation between the two related viruses let us conclude that the diverse coding strategies and hence different RNA structures play a role. All three genome segments of LACV and RVFV L and M segment are in negative polarity with only one transcriptional unit. RVFV S segment, however, uses ambisense coding strategy to direct synthesis of two proteins in an opposite direction, separated by a non-coding IGR (Giorgi et al., 1991, Elliott, 2014). By analyzing the predicted RVFV IGR, a hairpin structure can be observed with a central dsRNA stem with internal loops and peculiar poly-C/G repeats. These repeats were previously described as essential transcription termination signals (Albarino et al., 2007). As a minimal requirement for PKR activation serves a hairpin with a 16-base pair dsRNA stem flanked by 10 to 15 nucleotides of single-stranded tails (Zheng and Bevilacqua, 2004). The dsRNA stem thereby does not have to possess perfect base pairing since PKR can tolerate non-Watson-Crick structures and also internal loop structures within the dsRNA stem as long as the overall A-form geometry of the RNA is retained (Bevilacqua et al., 1998, Bevilacqua and Cech, 1996).

Thus, RVFV IGR presents all criteria of a potential PKR agonist. It was furthermore reported that PKR can interact with structured elements of viral RNA like HCV IRES or dimerized stem-loop structures of transactivation (TAR) RNA of HIV (Shimoike et al., 2009, Heinicke et al., 2009). However, whether PKR recognizes viral RNA packaged into nucleocapsids in infectious context remained elusive. Moy et al. previously reported in this regard that human DDX17 or Drosophila Rm62 specifically interacts with RVFV S segment IGR (Moy et al., 2014) indicating that the IGR would also be exposed for PKR interaction. PKR contacts the dsRNA mainly via 2′hydroxyl groups of the ribose sugar and oxygen residues of the phosphate backbone in a sequence independent manner (Bevilacqua and Cech, 1996).

Orthobunyavirus, like LACV, RNA is deeply sequestered within the nucleocapsids (Ariza et al., 2013) making the RNA highly inaccessible for PKR. RVFV nucleocapsids, however, are less symmetric and with few specific protein–protein interactions leaving the phosphate backbone partially exposed (Raymond et al., 2010, Raymond et al., 2012). Thus, oxygen groups of the phosphate backbone would be accessible for PKR interaction. Indeed, our data demonstrates that PKR is able to engage the IGR within
the viral nucleocapsid in the authentic context of virus infection. Interaction with the IGR promotes PKR conformational switching and phosphorylation identifying IGRs as natural PKR agonist. Partial exposure of the viral RNA alone, however, is not sufficient for PKR activation. This is indicated by the observation that RVFV M or S segmented nucleocapsids with the same packaging grade of the viral RNA only activate PKR if the IGR is present. As previously reported, influenza B virus nucleocapsids serve as PKR agonists at late stages of infection (Dauber et al., 2009). PKR, thereby, interacts with the 5’ppp dsRNA panhandle. Here, recognition of incoming nucleocapsids by PKR seems to be independent of the panhandle since RVFV M segmented nucleocapsids presenting a 5’ppp dsRNA panhandle failed to promote PKR activation. It can be postulated that at an early infectious stage, when low amounts of nucleocapsids are present RIG-I outcompetes PKR for 5’ppp panhandle interaction. It seems also plausible that PKR is only interacting with the 5’ppp promoter structure at later time-points of infection when misencapsidated side product of viral replication occur. Furthermore, PKR might not gain access to the panhandle of incoming nucleocapsids covered by the viral polymerase since PKR was not associated with the ability to remove proteins from RNA as it was postulated for RIG-I (Schmidt et al., 2011).

Interestingly, PKR interaction with the IGR of incoming nucleocapsids is independent on IFN induced upregulation of PKR as previously observed (Nallagatla et al., 2007). Already basal PKR levels are sufficient to recognize the IGR of viral nucleocapsids.

Despite RVFV, also representatives of the Arenaviridae use ambisense coding strategy, thus, bearing an IGR for transcriptional control. Arenavirus nucleoprotein sequesters the RNA bases like RVFV not completely leaving them partially exposed (Hastie et al., 2011b). The predicted IGR of TCRV, JUNV and LASV S and L segment represents a similar structure in comparison to RVFV with a hairpin composed of a dsRNA stem with internal loops and sporadic mismatches. Therefore, it was not surprising that like RVFV, also Old World Lassa virus and New World TCRV and JUNV nucleocapsids are able to promote PKR activation. This indicates, that the IGR presented in the nucleocapsids of two distinct virus families represent natural PKR agonist to promote an immediate antiviral response. Moy et al. postulated that interaction of a protein with the IGR of viral genomes has a negative impact on virus replication (Moy et al., 2014). We may, therefore, hypothesize that by interaction with bunyavirus RVFV and arenavirus TCRV, JUNV and LASSV IGR, PKR could also impair virus replication independent of its downstream signaling activity. However, this was beyond the scope of our study and requires further investigation.

PKR activation surprisingly could only be detected with incoming arenavirus nucleocapsids, but not if full infectious cycle was allowed. This raised the question whether arenaviruses encode for a PKR antagonist. By applying a diverse set of molecular biological assays LASV NP was identified as an inhibitor of PKR activation. Thereby, LASV NP interacts with PKR and induces its proteasomal degradation. It seems that the LASV NP encapsidating the viral genome does not affect the activation
status of PKR, which suggests that only free LASV NP acts as a PKR antagonist. LASV NP comprises dsRNA binding properties (Hastie et al., 2011a), hence, RNA could serve as a scaffold for PKR interaction. However, data supports rather a viral RNA independent interaction of LASV NP with PKR. Nevertheless, it cannot be excluded that LASV NP forms a complex with cellular RNA. It was reported in this regard that recombinant nucleoproteins of NSV can associate with cellular RNA into nucleocapsid-like structures (Ruigrok et al., 2011).

Interestingly, LASV NP comprises the ability to oligomerize (Lennartz et al., 2013). It remains to be resolved, whether NP monomers or LASV oligomeric complexes promote PKR antagonism. It was previously demonstrated that interference with the IFN response is not strictly dependent on NP oligomerization (Lennartz et al., 2013). Besides, Hastie et al. reported that rather the dsRNA specific 3’ to 5’ exonuclease activity of LASV NP is essential for inhibiting IFN signaling (Hastie et al., 2011a). Since incoming nucleocapsids do not present long dsRNA stretches to the host immune system, degradation of dsRNA might play a distinct immune evasion strategy during later infectious stages.

Arenavirus NP is well conserved between Old and New World arenaviruses (Hastie et al., 2011b), suggesting that an IFN antagonistic activity may be a shared feature (Hastie et al., 2011a). Indeed, Old World arenavirus LCMV and New World arenaviruses Whitewater Arroyo virus, Pichinde virus, JUNV, Machupo virus and Latino virus, but not TCRV were able to impair IFN and proinflammatory response (Martinez-Sobrido et al., 2006, Martinez-Sobrido et al., 2007, Rodrigo et al., 2012). However, the underlying mechanism still needs to be clarified. Likewise, arenaviruses use proteasomal degradation as a common PKR antagonism. However, the responsible viral protein could not be identified so far. TCRV unable to impair IFN signaling (Martinez-Sobrido et al., 2007) was surprisingly able to inhibit PKR activation. This highlights the requirement to prevent early recognition by PKR to ensure efficient virus propagation even in the case of apathogenic viruses like TCRV.

Conclusively, we identified the hairpin-structured IGR of RVFV S segment, New World arenavirus TCRV and JUNV and Old World arenavirus LASV nucleocapsids as natural PKR agonists. To prevent the antiviral PKR activity, TCRV, JUNV and LASV induce proteasomal degradation of activated PKR. In case of LASV, the newly synthesized nucleoprotein interacts with PKR and promotes its degradation via the proteasomal pathway. Hence, we identified the IGR as part of the viral nucleocapsid as an immediate PKR agonist in the authentic context of virus infection. Additionally, LASV NP was pinned down as a novel PKR antagonist. In addition to antagonizing immediate PKR recognition, arenavirus also impair early detection of viral nucleocapsids by RIG-I via an unpaired 5’ppp (Marq et al., 2010). Therefore, arenaviruses have evolved an efficient strategy to counteract immediate recognition and activation of antiviral defense mechanisms. Interestingly, IGRs play essential roles in the control of viral protein synthesis and hence virus replication (Lopez and Franze-Fernandez, 2007, Emery and Bishop, 1987, Geerts-Dimitriadou et al., 2012, Albarino et al., 2007), indicating the necessity to conserve this
structure despite the presence of antiviral defense mechanisms. This makes targeting of the IGR to block virus replication a promising target for drug development.
Acknowledgments

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Disclosures:

No conflicts of interest declared.
Figure Legend

Fig. 1: RIG-I and PKR differ in their agonist specificity
A549 cells were either left untreated (UT) or were treated with 50 µg/ml cycloheximide (CHX) and mock infected or infected with LACVΔNSs or RVFVΔNSs at an multiplicity of infection (MOI) of 5. After 5 h, cells were lysed and subjected to limited protease digestion (A and B), co-sedimentation (C) or co-immunoprecipitation assay (D). (A and B) Conformational switch. Cell lysates were treated with trypsin (+) and untreated cell lysates (-) served as input control. Western Blot analysis was performed with anti-RIG-I antibody (A) and with antibodies recognizing PKR and phosphorylated PKR (P-PKR) (B). (C) Co-sedimentation assay. Cell lysates were loaded on a 50-70% discontinuous glycerol gradient, centrifuged and collected fractions were concentrated. Samples were subjected to Western Blot analysis using antibodies directed against RIG-I, PKR and LACV or RVFV N as a marker for viral nucleocapsids. (D) Co-immunoprecipitation. Cell lysates were mixed with anti-p21 (negative control), anti-RIG-I or anti-PKR coupled magnetic beads and incubated for 2 h at 4°C. As input control 2% of the lysates were kept. Input and eluates were analyzed by Western Blotting using anti-RIG-I, anti-PKR, anti-LACV or RVFV N and anti-p21 antibodies.

Fig. 2: Intergenic region of incoming RVFV nucleocapsids serve as PKR agonists.
(A) RVFV intergenic region. Secondary structure prediction of RVFV S segment (data base entry EU312104.1) intergenic region via free energy minimization (mFold). (B) RVFV coding strategy. Viral particles contain three genome segments in negative polarity (vRNA) which are named large (L), middle (M) and small (S). L and M segment contain only a single transcriptional unit whereas the S segment uses ambisense coding strategy. Thereby, two open reading frames are separated by an intergenic region (IGR). During replication, each genome segment is first transcribed to allow protein synthesis of essential proteins for genome replication. Amplification of the viral genome (vRNA) involves the synthesis of an exact copy of the vRNA, called complementary RNA (cRNA). In case of the S segment, the cRNA serves also as a template for mRNA synthesis. (C) Conformational switch. A549 cells were treated with 50 µg/ml CHX and then infected for 5 h with RVF VLPs containing either M or S segmented nucleocapsids. Limited protease digestion and analysis of the samples was performed as for Fig. 1A and B. (D) Super-resolution Immunofluorescence Microscopy of RIG-I and PKR nucleocapsid complexes. Cells were treated and infected as described for C and then fixed and stained against RVFV N (green channel) and RIG-I (upper panel; red channel) or PKR (lower panel; red channel). Quantification was performed of three independent experiments. Student’s T-test was carried out for statistical analysis. Significant differences are indicated by asterisks (** p<0.01).
Fig. 3: Arenavirus replication products prevent nucleocapsid promoted PKR activation.

(A) Secondary structure prediction of TCRV S segment (data base entry NC_004293.1), TCRV L segment (data base entry NC_004292.1), JUNV S segment (data base entry JF799984.1), JUNV L segment (data base entry JF799980.1), LASV S segment (data base entry HQ688673.1) and LASV L segment (data base entry HQ688674.1) intergenic region via free energy minimization (mFold). (B) PKR activation status. A549 cells were either left untreated (UT) or were treated with 50 µg/ml CHX and mock infected or infected with RVFVΔNSs, TCRV, JUNV or LASV at an MOI of 1. Infection was stopped 5 h post-infection and cell lysates were prepared with 0.5% Triton X-100 in PBS. Limited protease digestion and analysis of the samples was performed as for Fig. 1B.

Fig. 4: LASV nucleoprotein promotes proteasomal degradation of PKR.

(A) Interaction of RIG-I and PKR with LASV nucleocapsids. A549 cells were left untreated (UT) or were treated with 50 µg/ml CHX and either mock or LASV infected at an MOI of 5. After 5 h, cell lysates were prepared and co-immunoprecipitation was performed as described for Fig. 1D. For Western Blot analysis anti-RIG, anti-PKR, anti-P-PKR, anti-LASV NP and anti-p21 antibodies were used. (B) Overexpression analysis. HEK293T cells were transfected with either 0.25, 0.5, 1 or 2 µg of HA-tagged expression constructs for LASV NP, GP or Z together with 1 µg PKR plasmid. Normalization of transfected plasmid amount was done with ΔMx plasmid. Expression was allowed for 24 h and then cell lysates were prepared. Samples were analyzed by Western Blotting using anti-P-PKR, anti-PKR and anti-HA antibodies. (C) Co-immunoprecipitation. HEK293T cells were transfected with 1 µg of LASV NP, GP or Z together with 1 µg PKR plasmid. Lysates were prepared 24 h post-transfection and subjected to co-immunoprecipitation with anti-PKR coupled beads as described in Fig. 1D. Input samples and eluates were validated as described for B. (D and E) Influence of proteasomal degradation. (D) HEK293T cells were transfected like C and 8 h post-transfection either left untreated or treated with 20 µM MG132. After additional 16 h, cells were lysed and analyzed as described for B. (E) Conformational switch. A549 cells were infected with RVFVΔNSs, TCRV, JUNV or LASV at an MOI of 1 and after the inoculum was removed the cells were left either untreated (UT) or were treated with 20 µM MG132. After 5 h, cells were lysed and limited protease digestion and Western Blotting was performed as described for Fig. 1B.

Fig. 5: Mechanism of LASV NP mediated PKR antagonism

Arenavirus IGR exposed from the viral nucleocapsid can be recognized by PKR. Once activated, PKR blocks cap-dependent translation of viral and host cell proteins by phosphorylation of the alpha subunit of eukaryotic initiation factor 2 (eIF2α). Additionally, PKR induces an antiviral type I interferon
(IFN) and proinflammatory response. However, to circumvent PKR activation, LASV NP interacts with PKR and promotes its proteasomal degradation.
Supplemental information

Supplemental Experimental Procedure

Co-sedimentation assay of recombinant LCV and RVFV nucleoprotein

For co-sedimentation assays, A549 cells grown in two wells of a 6 well plate were transiently transfected with 1 µg pl.18_RVFV N or pTM_LACV N using Nanofectin transfection reagent (PAA Laboratories). Cell lysates were prepared 24 h post-transfection and subjected to co-sedimentation assay as ascribed above. Western Blot analysis was performed with mouse monoclonal anti-RIG-I antibody at 1:1000, mouse monoclonal anti-PKR B10 at 1:500, rabbit polyclonal anti-LACV N at 1:1000, and mouse polyclonal anti-RVFV N serum at 1:1000.

Time-course experiment

A549 cells were infected with LACVΔNSs, RVFVΔNSs, TCRV, JUNV or LASV at an MOI of 1. If indicated, cells were treated with 50 µg/ml CHX 1 h prior, during infection and while the course of the experiment. After 2, 5 and 24 h cells were washed and lysed in 1-fold sample buffer. Samples were analyzed by SDS-PAGE and Western Blotting using mouse monoclonal anti-PKR B10 at 1:500, rabbit monoclonal anti-p-PKR Thr 446 at 1:500, rabbit polyclonal anti-LACV N at 1:1000, mouse polyclonal anti-RVFV N serum at 1:1000, rabbit monoclonal anti-LASV NP at 1:3000, guinea pig polyclonal anti-JUNV NP at 1:500 and guinea pig polyclonal anti-TCRV NP at 1:500. Staining with mouse monoclonal anti-actin served as a loading control.

Suppl. 1: RIG-I and PKR do not interact with LACV and RVFV nucleoprotein

Co-sedimentation assay. HEK293T cells were transfected with either LACV or RVFV N and 24 h post-transfection cell lysates were prepared. Samples were loaded on a 50-70% discontinuous glycerol gradient, centrifuged and collected fractions were concentrated. Samples were subjected to Western Blot analysis using antibodies directed against RIG-I, PKR and LACV or RVFV N as a marker for viral nucleocapsids.

Suppl. 2: PKR activation during bunyavirus and arenavirus infection

PKR activation status. A549 cells were either left untreated (UT) or were treated with 50 µg/ml CHX and mock infected or infected with LACVΔNSs, RVFVΔNSs, TCRV, JUNV or LASV at an MOI of 1. After 2, 5 and 24 h cell lysates were prepared with 1-fold sample buffer and samples were analyzed by Western blotting using antisera against phosphorylated PKR, PKR and LACV N, RVFV N, TCRV NP, JUNV NP or LASV NP, respectively. Staining against actin served as a loading control.
References


21


nucleocapsids containing a 5'-triphosphorylated genome activate RIG-I and antiviral signaling. Cell Host Microbe, 13, 336-46.


### Figure 4

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- **LASV**
  - RIG-I
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  - P-PKR
  - NP
  - p21

#### 4B

- **LASV NP-HA**
- **LASV GP-HA**
- **LASV Z-HA**

- P-PKR
- PKR
- HA

#### 4C

- LAVS Z-HA
- LASV GP-HA
- LASV NP-HA
- delIMX

- PKR
- P-PKR
- HA

#### 4D

- UT
- MG132

- LASV Z-HA
- LASV GP-HA
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- delIMX

- PKR
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- NP
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#### 4E

- trypsin
- mock
- RVFV
- TCRV
- JUNV
- LASV

- PKR
- P-PKR

- PKR resistant fragment

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- CHX
- MG132
Suppl. 1

50% glycerol  RVFV N  70% glycerol

1 2 3 4 5 6 7 8 9 10 11 12

RIG-I
PKR
RVFV N

50% glycerol  LACV N  70% glycerol

1 2 3 4 5 6 7 8 9 10 11 12

RIG-I
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4.5. REVIEW: RIG-I-like receptors and negative-strand RNA viruses: RLRly bird catches some worms

Own contribution:

I contributed to writing of the manuscript.

____________________________
Michaela Gerlach
Mini review

RIG-I-like receptors and negative-strand RNA viruses: RLRly bird catches some worms

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A R T I C L E   I N F O

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Keywords:
Negative-strand RNA virus
Genome segmentation
RIG-I-like receptors
Panhandle
Viral countermeasures

A B S T R A C T

Negative strand RNA viruses with a nonsegmented genome (ns-NSVs) or a segmented genome (s-NSVs) are an important source of human and animal diseases. Survival of the host from those infections is critically dependent on rapidly reacting innate immune responses. Two cytoplasmic RNA helicases, RIG-I and MDA5 (collectively termed RIG-I-like receptors, RLRs), are essential for recognizing virus-specific RNA structures to initiate a signalling cascade, resulting in the production of the antiviral type I interferons. Here, we will review the current knowledge and views on RLR agonists, RLR signalling, and the wide variety of countermeasures ns-NSVs and s-NSVs have evolved. Specific aspects include the consequences of genome segmentation for RLR activation and a discussion on the physiological ligands of RLRs.

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1. Introduction

Viruses with a single-stranded (ss) RNA genome of negative polarity (negative-strand RNA viruses, NSVs) are responsible for a wide range of diseases. Members of the NSV group cause respiratory problems (e.g., influenza, hantavirus pulmonary syndrome, respiratory syncytial virus), severe childhood diseases (measles, mumps), neuronal infection (La Crosse virus, Toscana virus, Borna disease virus), even with up to 100% mortality (Rabies), as well as multi-organ failure and haemorrhagic fevers (Rift Valley fever, Severe fever with Thrombocytopenia, Nipah, Ebola, Lassa, Hanta, Crimean Congo haemorrhagic fever).

NSVs encode highly active polymerases, producing in a typical case 100,000 RNA copies per cell within 10 h of infection [1]. Quite expectedly, such numbers of non-self RNAs in the cytoplasm are highly immunogenic and could trigger massive antiviral responses. To avoid being defeated by the innate immune system, NSVs have evolved strategies against recognition, antiviral signalling, and the launched antiviral mechanisms. In this review article, we will attempt to provide an update on the principles of cytoplasmic virus recognition and viral escape. For deeper insights into specific aspects and into the extracellular virus recognition which is not covered here, we refer to the wealth of recent, excellent reviews by others [2–14].

2. Negative-strand RNA viruses

Taxonomically, the group of NSVs is divided into those having one, continuous strand of genomic RNA (the Mononegavirales) and those whose genome is divided into several segments. The Mononegavirales, also called nonsegmented NSVs (ns-NSVs) have all genes lined up along the ssRNA genome, separated by regulatory intergenic regions acting as transcriptional promoters, and flanked by the regulatory leader and trailer sequences (Fig. 1A). For segmented NSVs (s-NSVs), by contrast, each RNA segment contains one promoter (formed by annealing of the 5‘ and 3‘ ends) driving expression of one gene (Fig. 1B). The number of segments differs between s-NSV families. Members of the Orthomyxoviridae possess mostly 8 segments, whereas Bunyaviridae have 3 and Arenaviridae have 2 segments (Table 1). Some bunyaviruses and all arenaviruses enlarge the genetic capacity of their segments by transcribing an additional gene from the copy of the genome RNA (the cRNA), the so-called ambisense strategy.

3. Intracellular virus recognition by RIG-I like receptors

A rapid and efficient antiviral response is essential to limit virus replication and ensure survival of the host. The cytoplasmic RIG-I-like receptors (RLRs) are able to detect viral RNA species and to trigger an intracellular signalling cascade that eventually results in the release of specific cytokines and chemokines [2,10]. Type I interferons (IFN-α/β) represent the most important antiviral cytokines, inducing expression of more than 300 IFN-stimulated
genes (ISGs) [12]. ISGs act in a variety of ways to inhibit virus replication, to elevate antiviral alertness in the infected and surrounding cells, and to modulate the adaptive immune system.

3.1. RLR structure

The group of RLRs consists of RIG-I (retinoic acid inducible gene 1), MDA5 (melanoma differentiation association factor 5) and LGP2 (laboratory of genetics and physiology 2) [10]. RLRs are members of the DExD/H box RNA helicase family and consist of a carboxy-terminal domain (CTD), a central DExD/H box RNA helicase domain (composed of Hel1, Hel2 with the insertion domain Hel2i in between) and, in the case of RIG-I and MDA5, two additional amino-terminal caspase recruitment domains (CARDs) (Fig. 2) [15,16]. For RIG-I, the CTD and the helicase domain are conducting ligand recognition, whereas in case of MDA5 this function seems to be performed by the helicase domain. The terminal CARDs are required for signal transduction. LGP2 does not signal by itself but can bind double-stranded RNA (dsRNA) and acts as a positive regulator of MDA5 [17,18].

3.2. RLR signalling

RIG-I has in non-infected cells an auto-repressed conformation in which the CARDs are masking the RNA binding site of the helicase domain [16]. The CTD is exposed to the cytosol and connected by a flexible linker. When the CTD binds to a specific ligand, RIG-I switches its conformation to wrap around the RNA ligand (via CTD and helicase domain) and expose the CARDs instead of the CTD [9]. Stabilization of the active conformation and homo-oligomerization of RIG-I are facilitated by post-translational modifications like dephosphorylation, covalent K63 ubiquitylation via TRIM25, and association of unanchored ubiquitin chains [4,19–23]. Activated RIG-I can form helical tetramers, but – similar to MDA5 – also longer oligomers along dsRNA [21,24,25].

The oligomers of RIG-I and MDA5 serve as platforms to recruit the adaptor protein MAVS (mitochondrial antiviral signalling) via multiple CARD–CARD interactions [2]. Activated MAVS associates with TRAF3/6 to promote activation of the kinases TBK1/IKKε which are responsible for the phosphorylation of the transcription factors IRF3 and IRF7 [2]. In parallel, the kinases IKKα and IKKβ can activate NF-κB. IRF3/7 and NF-κB translocate into the nucleus to initiate expression of IFN-α/β and proinflammatory cytokines. The basics of RLR signalling are outlined in Fig. 3 (middle panel).

Fig. 1. Differing coding strategies of nonsegmented and segmented NSVs. Symbolized virus particles are shown on top, followed by a sketch of the unencapsidated viral genome with the individual genes as grey boxes. By convention the ssRNA genomes of NSVs are drawn from 3′→5′. (A) The genome of ns-NSVs consists of one continuous stretch of ssRNA from which all mRNAs are synthesized. Intergenic regions contain stop/start signals regulating termination and initiation of transcription. Note the uncapped 5′ppp RNA transcribed from the leader region. Basic set-up of a Rhabdovirus is shown as example. (B) s-NSVs transcribe mostly one mRNA per genome segment (with the exception of ambisense segments which express another gene transcribed from the opposite direction). n = number of segments. Not all s-NSVs have mRNAs with a polyA, which is therefore shown in brackets.

![Fig. 1](image-url)

<table>
<thead>
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<th>Table 1 Negative-strand RNA viruses.</th>
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<tr>
<td><strong>Family</strong></td>
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<td>Bunyaviridae</td>
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<td>Orthomyxoviridae</td>
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*Virus acronyms and genera shown in brackets

![Fig. 2](image-url)
3.3. RLR agonists

Despite their structural and functional similarity, RIG-I and MDA5 recognize different RNA ligands (see below), and are activated by distinct but overlapping subsets of viruses [13]. Ligand-wise, RIG-I is strongly activated by blunt-ended dsRNAs of at least 10 base pairs length bearing a 5′triphosphate (5′ppp) group [15,26,27], by long dsRNA molecules of more than 200 base pairs (irrespective of the 5′ end) [24,28], but also by 3′-monophosphorylated ssRNAs [29] and poly U/UC rich stretches of ssRNA [30]. MDA5 detects long dsRNA molecules, ideally with higher order RNA structures [31], and can be activated by particular stretches of NSV RNAs [32,33]. Both RLRs also show preference for AU-rich ssRNAs which seems independent of structure [33,34].

While these activating molecules had mostly been identified and characterized by transfections and manipulations of naked RNAs, they nonetheless allowed conclusions on the nature of the physiological RLR agonists present in infected cells (summarized in Fig. 3, left and right panels). NSVs encapsidate their ssRNA genome and its template (and copy), the antigenome, with nucleocapsid proteins [35]. Therefore, NSVs do not produce detectable amounts of RLR-critical dsRNA [36]. However, synthesis of the NSV genome is initiated with an unprimed nucleoside triphosphate, resulting in a 5′ppp terminus. For ns-NSVs, nucleocapsids are linear [35], suggesting that the 5′ppp ssRNA end does not meet the 3′ counterpart and hence cannot activate RIG-I [13]. For s-NSVs, by contrast, the promoter structure which is formed by annealing of the genome termini ("panhandle"; see Fig. 1B) bears hallmarks of a RIG-I agonist. Indeed, we recently showed that the 5′ppp dsRNA panhandle structures packaged into nucleocapsids of two members of the Bunyaviridae (La Crosse virus (LACV) and Rift Valley Fever virus (RVFV)) act as RIG-I agonists in the natural context of infection [37]. Therefore, RIG-I can recognize the nucleocapsids of s-NSVs directly after their entry into the cytoplasm. In the case of ns-NSVs, only RNA synthesis products (e.g. 5′ppp leader RNA) can act as triggers of RIG-I [6,20,33,38–41]. Moreover, particular stretches of viral mRNA can activate MDA5 [32,33]. During replication of the RNA genome, the viral polymerase occasionally jumps template, resulting in RNA products with internal deletions, the defective interfering (DI) RNAs, named so because they strongly compete with the full-length genomes for resources [42]. One particular class, the copy-back DI RNAs, contain complementary 5′ and 3′ ends. It is thought that a percentage of DI RNAs is incompletely encapsidated [43]. For the ns-NSV Sendai virus (SeV), it was shown that copyback DI RNAs form panhandle-like dsRNA structures which can accommodate RIG-I oligomers and activate IFN induction [20,43]. Also for s-NSVs, products of viral RNA synthesis (including internally deleted genomes) can activate RIG-I and innate immune signalling [34,44–46]. Thus, taken together, for s-NSVs incoming nucleocapsids can be detected by RIG-I, and also later RNA synthesis products trigger innate immunity. For ns-NSVs, by contrast, the RLRs have to await the appearance of regular and irregular products of viral RNA synthesis.

4. Pros and cons of genome segmentation

s-NSVs are schlepping a bundle of genome segments, each with a RIG-I-sensitive 5′ppp dsRNA promoter. This clearly represents a
disadvantage with respect of immune activation. Moreover, segmentation requires sophisticated packaging mechanisms to ensure each virion contains the full set of genes. On the other hand, the segments can easily be exchanged between different virus variants (reassortment), allowing rapid emergence of new viral strains with altered antigenic or replicatory properties [5,47]. Dividing the genome in separate units is also considered an insurance against sequence degeneration in systems with high error rates, such as RNA-dependent polymerases [48], and packaging several shorter segments instead of one large genome was recently shown to increase the physical stability of virus particles [49]. One evolutionary trade-off of genome segmentation, however, is the elevated number of RLR agonists presented to the innate immune system. It is therefore conceivable that s-NSVs have to express early-hitting IFN evasion strategies which may also be stronger than those of ns-NSVs.

5. RLR evasion strategies of NSVs

To our knowledge, all NSVs investigated so far have evolved countermechanisms of RLR signalling, acting to prevent recognition by RLRs, to interfere with RLR signalling, and/or to suppress

<table>
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<tr>
<th>Interference with</th>
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<th>Virus</th>
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<td>VP35</td>
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<td>Recruitment repression factor SAP30 to inhibit IFN-β transcription</td>
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<td>Drives proteasomal degradation of RPB1</td>
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induction of the type I IFN response. The molecular mechanisms involved range from a selective interference with key components of the IFN system to a broad shut-off of the host cell transcription. Since it has become virtually impossible to summarize all known mechanisms and factors of viral IFN antagonism, we can only describe prominent examples to highlight the underlying principles (Table 2).

5.1. Prevention of RLR sensing

An efficient way of innate immune escape is to avoid the initiation of RLR signalling directly at its root, i.e., at the level of detection. As mentioned, NSVs do normally not produce long dsRNAs [36], first of all because of the packaging of genome and antigenome RNAs into separate nucleocapsids [35]. For the s-NSV influenza A virus (FLUAV), depletion of the cellular helicases UAP56 and URH49, which are known to be involved in RNA encapsidation [50], resulted in the appearance of long dsRNA [51]. Also for an ns-NSV (vesicular stomatitis virus, VSV) UAP56 and URH49 are necessary for prevention of dsRNA formation, suggesting a general mechanism [51]. In addition, ns-NSVs are recruiting the cellular RNA binding protein La to shield the leader RNA from RIG-I recognition [39]. The generation of immuno-active RNAs products can also be avoided by tightly controlling the processivity of genome transcription and replication. For several paramyxoviruses (i.e. ns-NSVs) it was described that a loss of regulatory proteins (dependent on the virus either P, C, V) or a promotor mutation enhanced viral RNA synthesis but also facilitated appearance of innate immunity-inducing dsRNA, DI particles, or other aberrant RNAs [52–57]. A similar control of RNA synthesis and DI particle formation to avoid IFN induction was recently reported for FLUAV, suggesting that this is a general mechanism of NSVs [45].

RLR sensing could also be a reason why members of the Orthomyxoviridae are replicating in the nucleus. This virus family contain 6 (Thogoto virus, THOV) to 8 (FLUAV) genome segments, i.e. the virions contain at least 2 times as many RLR ligands as those of other s-NSVs. Since RLRs are localized in the cytoplasm, they are only transiently in contact with orthomyxoviral nucleocapsids, namely during the initial transport from the endosome to the nucleus, and during the late step of particle egress. Hiding in the nucleus could be thus a way by which heavily-segmented NSVs can minimize RLR sensing. Indeed, innate immunity activation by FLUAV requires RNA synthesis and RNA nuclear export [44,58]. Thus, the early cytoplasmic passage of nucleocapsids can occur largely unnoticed by the cell.

On top of the strategies to prevent production or exposure of RLR-relevant RNAs, both ns-NSVs and s-NSVs exhibit a range of active mechanisms. The proteins VP35 of Ebola (EBOV) and Marburg virus (MARV) and NS1 of FLUAV directly bind dsRNA [59–62], although it seems that additional host factor interactions are necessary to prevent IFN induction [63–66]. Moreover, members of the Bornaviridae (ns-NSVs) and Bunyaviridae (s-NSVs) remove the 5’ppp group from their genome to avoid RIG-I activation [67,68], although some residual activating potential seems to remain [37,69]. Similarly, for the Arenaviridae (s-NSVs) a fraction of genome ends [70] can exhibit a single 5’overhanging nucleotide with the potential to disturb RIG-I function [71]. Moreover, Lassa virus (LASV, family Arenaviridae) encodes a nucleoprotein with 3’–5’ exoribonuclease activity to degrade dsRNA [72,73].

5.2. Specific interference with RLR signalling

In addition to masking, processing or degrading RLR agonists, NSVs can interfere with key components of the RLR signalling pathway. Several factors of ns-NSVs and s-NSVs are known to act on the level of RLRS. NS2 of respiratory syncytial virus (RSV) and the Z protein of New World Arenaviruses directly interact with RIG-I to prevent the association with MAVS [74,75]. The ovarian tumour (OTU) domain of Nairoviruses (family Bunyaviridae) suppresses IFN induction and de-ubiquitylates RIG-I [76,77]. MDA5 was originally discovered as a host interactor and inhibitor of the paramyxovirus IFN antagonist V [78]. The V proteins are wedging into the MDA5 structure in a manner that prevents assembly into signalling-competent filaments [79]. VP35 of EBOV sequesters the RIG-I activating protein PACT to interfere with RIG-I activation [64]. V proteins of paramyxoviruses assemble RIG-I and LGP2 into a complex which becomes refractory to activation by RIG-I ligands [80], and NS1 of FLUAV interacts with TRIM25 to counteract ubiquitinylination of RIG-I [63]. Interestingly, infections with VSV or SeV were shown to upregulate the lectin Siglec-G, resulting in RIG-I degradation via K48-linked ubiquitinylination [81]. A strategy of ubiquitin-dependent RIG-I degradation is also conducted by Toscana virus (TOSV) via its NSs protein [82].

Some viral factors also sequester key components of RLR signalling in cellular compartments. RSV N and P proteins recruit MDA5, MAVS, and less efficiently RIG-I, into virus induced inclusion bodies [83]. Also Severe fever with thrombocytopenia syndrome virus (SFTSV) NS5 prevents RLR signalling by promoting relocation of RIG-I, TRIM25, TBK1, IKKε and IRF3 into virus-induced cytoplasmic structures [84,85]. NS1 and NS2 of RSV induced the formation of the so-called degradosome to target MAVS and IRFs for degradation [86]. MAVS signalling is impaired by FLUAV polymerase subunits (PB1, PB2 and PA) [87,88] and the accessory protein PB1-F2 [89–91]. Further downstream, TBK1 activation or action are prevented by N of arenaviruses and hantaviruses, and Gn proteins of hantaviruses, P proteins of BDV and Babies virus, V proteins of paramyxoviruses, and also by EBOV VP35 [66,92–97].

Finally, at the level of IFN transcription, several viral factors are known to interfere with IRF or NF-kB activity. EBOV VP35 (a multifunctional IFN antagonist like FLUAV NS1) enhances SUMOylation of IRF7 via the cellular E3 ligase PIAS1, thereby inhibiting its function as an IFN transcription factor [98]. ML of THOV (family Orthomyxoviridae), acts by blocking IRF3 and IRF7 signalling and association with CBP, TRAF6 and the general transcription factor IIB [99,100]. V proteins of several (but not all) paramyxoviruses interact with IRF3 and impair its nuclear translocation [101]. In case of the highly pathogenic Nipah virus (NIV) (family Paramyxoviridae), the W protein translates into the nucleus to inhibit activation of the IFN-β promoter [102]. Also the NSs protein of RVFV specifically inhibits IFN-β mRNA transcription by forming a promoter-bound complex with the repression factor SAP30 [103].

5.3. Unspecific interference with RLR signalling

In contrast to the above-described specific mechanisms aiming at antiviral signalling pathways, some representatives of the NSVs interfere with the cellular transcription machinery in total. It is thereby important to distinguish between a primary host cell shut-off imposed by specific viral mechanisms from the secondary shut-off caused by translation inhibition due to the dsRNA-activated host cell kinase PKR. While the primary, “intended” shut-off seems to be mostly (but not exclusively) the domain of s-NSVs, the secondary shut-off is in fact an efficient antiviral host reaction. PKR activation is thus avoided by both s-NSVs and sn-NSVs, either by tight regulation of RNA synthesis (see above), or by specific anti-PKR mechanisms [31,104].

A virus-induced, general block of gene expression provides an efficient mechanism to prevent synthesis of IFN and ISGs, but could
bear the disadvantage of cutting off from renewable resources. However, viruses undergoing rapid and lytic infection cycles seem to care little for such considerations. That the viral host shut-off, a long-known phenomenon, is often an IFN antagonism in disguise was revealed by genetic complementation studies showing that the particular viral shut-off factor is only necessary if the host organism possesses a functional IFN system [105–107]. In other words, in an IFN-free host environment the shut-off factors are dispensable for viral success. Interestingly, all s-NVs (but not ns-NSVs) commit cap-snatching, an endonuclease-mediated mechanism to degrade host mRNAs by depriving them of their 5′ cap structure. Cap-snatching spares the s-NVs to encode their own mRNA capping enzyme, and could have the convenient side effect of reducing host cell gene expression, including that of upregulated components of the RLR and IFN signalling pathways [108]. FLUAV expresses its endonuclease even as a separate domain, called PA-X, which was actually shown to suppress antiviral host cell responses [109]. FLUAV impairs host gene expression also by other mechanisms. The multifunctional protein NS1 interferes with processing, nuclear export and translation of host mRNAs [8]. Inhibition of nuclear-cytoplasmic RNA transport is a strategy shared by the matrix protein M of VSV [110]. Broad disruption of RNA polymerase II activity is achieved by representatives of the Bunyaviridae. The NSs of RRVF disturbs assembly of the general transcription factor TFIIH [111] and promotes degradation of the TFIIH subunit p62 via the E3 ubiquitin ligase FBXO3 [112,113]. The NSs protein of Bunyamwera virus and LACV impact on the RNA polymerase II subunit RB1, disturbing its phosphorylation (and hence processivity) or driving its proteasomal degradation [114,115].

6. Conclusions and outlook
NSV-infected cells are flooded with foreign RNAs which are – compared to the pool of host cell RNAs – rather homogenous in sequence and structure. Unusual features like double-strandedness or triphosphorylated 5′ ends are extremely efficient markers of infection. However, also structures with less prominent features are conceivable to act as triggers of antiviral responses if they appear in high enough amounts.

Since the discoveries of RIG-1 and MDA5 [78,116], the research field of intracellular RNA recognition and viral countermechanisms has virtually exploded. There are ongoing discussions on the physiological ligands of RLRs and on major RLRs engaged during a particular infection, questions which are touching the very core of the immune response. The distinction between self and non-self. In our review, we wanted to highlight that NSVs are a quite heterogeneous taxonomic group, with different genome and nucleocapsid structures, subcellular localizations, replication strategies, tendencies to produce DI particles, and a multitude of IFN antagonists which can obscure RLR-ligand interactions. Moreover, different infection phases may produce different RLR triggers. To make things even more complicated, it is meanwhile established that RLRs can get help from accessory factors such as e.g. other RNA-binding proteins, helicases or RNases, further expanding the range of in vivo RLR triggers [17,32,39,117–119], which may not work as such in vitro. So, again, the question which RLR recognizes which virus and viral structure under physiological conditions may return different answers which depend on the infection phase, the DI particle content of virus stocks, the specific activity of virus regulators and IFN antagonists, and on host cell cofactors.

Conflict of interest
No conflicts of interest declared.

Acknowledgments

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4.6. REVIEW: Segmented negative-strand RNA viruses and RIG-I: divide (your genome) and rule

Own contribution:

I contributed to writing of the manuscript.

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Michaela Gerlach
Segmented negative-strand RNA viruses and RIG-I: divide (your genome) and rule
Michaela Weber and Friedemann Weber

The group of negative-stranded RNA viruses (NSVs) with a segmented genome comprises pathogens like influenza virus (eight segments), Rift Valley fever virus and Hantavirus (three segments), or Lassa virus (two segments). Partitioning the genome allows rapid evolution of new strains by reassortment. Each segment carries a short double-stranded (ds) ‘panhandle’ structure which serves as promoter. Similar dsRNA structures, however, represent the optimal ligand for RIG-I, a cytoplasmic pathogen sensor of the antiviral interferon response. Thus, segmenting a virus genome can entail an increased RIG-I sensitivity. Here, we outline the astonishingly diverse and efficient strategies by which segmented NSVs are compensating for the elevated number of RIG-I ligands in their genome.

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Edited by Maria-Carla Saleh
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Introduction
Like eukaryotes, some negative-strand RNA viruses (NSVs) feature a genome which is divided into semi-autonomous pieces. The evolutionary advantage is an extra degree of freedom for recombination, facilitating the rapid admixture of genome segments to create hybrid offspring. The success of this strategy is obvious from the constant flow of new influenza strains afflicting mankind with epidemics or even pandemics [1]. Besides influenza viruses (belonging to the family Orthomyxoviridae) which have up to eight segments to offer for genetic mixing, there are two other families with a more modest set-up, namely the Bunyaviridae (three segments) and the Arenaviridae (two segments). Together these viruses are responsible for an impressive array of diseases ranging from severe pulmonary, hepatic or renal ailments (influenza, Rift Valley fever, Hantavirus) up to deadly hemorrhagic fevers (Lassa fever, Junin, Crimean Congo hemorrhagic fever, Hantavirus, Rift Valley fever).

Host organisms (e.g., us) are however not defenceless. Adaptive immunity would be too slow to protect from a first infectious encounter, but mammals can execute an immediate innate immune response enabled by genome-encoded factors specialized to rapid virus detection. A well-established example is given by the cytoplasmic pathogen recognition receptor RIG-I, an RNA helicase of the DExD/H type [2]. One of the best ligands for RIG-I is short double-stranded RNA (dsRNA) containing a blunt 5’ end with a triphosphate group, 5′ ppp dsRNA [3] (Figure 1a). Ligand-activated RIG-I is rapidly triggering a signalling chain which upregulates an antiviral innate immune response. 5′ ppp dsRNA structures similar to the ideal RIG-I ligand are formed by the genome ends of segmented NSVs (Figure 1b), but are not found on cellular RNA [3]. Thus, RIG-I seems well adapted to the genome of segmented NSVs, ensuring a specific and rapid immune reaction to this class of viruses.

In this opinion article, we will briefly outline how RIG-I-dependent ligand recognition and antiviral signalling work, how segmented NSVs are activating it, and share our thoughts on how these viruses are trying to escape detection and signalling by RIG-I.

RIG-I-mediated antiviral response
RIG-I is composed of an amino-terminal tandem of CARDs (caspase recruitment domains), a central RNA helicase domain, and a carboxy-terminal CTD (C-terminal domain) [4]. Ligand-mediated activation of RIG-I results in the binding of the adaptor protein MAVS (mitochondrial antiviral signalling) via polyubiquitin-mediated CARD–CARD interactions. After recruitment of further cofactors, the complex activates the transcription factors IRF-3/7 and NF-κB, which, in turn, initiate transcriptional activation of type I interferons (IFN-alpha/beta) and other proinflammatory cytokines [5]. Once secreted, IFNs bind to their cognate receptor on the cell surface to activate numerous ISGs (IFN-stimulated genes). Many ISGs encode antiviral restriction factors inhibiting different steps of virus replication such as entry, transcription, replication, translation, assembly or release from infected cells, thus slowing down or entirely blocking further viral spread [1,4–6].
Activation of RIG-I by viral structures

In uninfected cells, RIG-I is present in an auto-inhibited conformation stabilized by internal interactions between the inner CARD (CARD2) and a part of the helicase domain, the Hel2i region [7,8,9**]. The CTD is flexibly exposed and thus available for scanning the cytosol for viral RNAs. Binding of 5’ ppG dsRNA to the CTD induces a massive conformational switch by disrupting

Synthetic and natural RNA structures relevant for RIG-I signalling. (a) Minimal RIG-I agonist as determined by RNA transfection experiments [7,10]. (b) Idealized structure of segmented negative-sense RNA genomes, with a large single-stranded coding region pseudocircularized by a double-stranded ‘panhandle’ promoter. (c)-(f) Predicted structures of viral panhandle regions. (c) Influenza A virus (Orthomyxoviridae) segment 8 [16,22]. (d) Rift Valley fever virus segments M (database entry DQ380206) and L (database entry DQ375403). (e) Hantaan virus segments S (database entry AF288644) and M (database entry KC344261). Note the monophosphate at the 5’ end. (f) Junin virus segments S (database entry NC_005080) and L (database entry NC_005081), both as corrected by Albaríño et al. [24]. The unpaired nucleotide at the 5’ end found in a fraction of the S segment RNAs [24] is shown in lower case. In all subfigures, Watson-Crick basepairs are indicated by lines, and non-Watson-Crick basepairs [50] are indicated by dots. Sequences are in the viral sense orientation and nucleotides are numbered up to the first interruption of base pairing.

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In the Hel2i–CARD2 interaction, thereby clamping RIG-I around the RNA ligand and liberating the CARDs to interact with polyubiquitin and MAVS [4,6].

Besides 5′ppp dsRNA of a minimal length of 10 bp [7,10], RIG-I responds to long dsRNA molecules of more than 200 bp (irrespective of the 5′ ends), to 3′-phosphorylated cleavage products of RNase L, and to poly-U/UC-rich ssRNA stretches [3,11–13]. Long dsRNA is not produced in a detectable amount during the course of NSV infection [14]. However, as outlined above, segmented NSVs form a RIG-I-critical 5′ppp dsRNA region by base-pairing the terminal 5′ and 3′ non-translated regions of their genome (Figure 1c–f). This so-called ‘panhandle’ structure pseudo-circularizes the otherwise single-stranded RNA genome and serves as promoter for transcription and replication [15–17]. Moreover, at least for influenza viruses a 3′ untranslated ssRNA region of the genome activates RIG-I in a 5′ppp independent fashion [18*].

In most of the RIG-I studies mentioned above, the structural characteristics of the ligands had been determined in vitro or by transfection of naked RNA into cells. Hence, the question was raised which of those structures are actually representing the natural ligands of RIG-I. Rehwinkel et al. [19] and Baum et al. [20] showed that full-length virus genomes and defective shortened RNAs arising during the course of the infection are indeed serving as RIG-I activators. Moreover, our group recently demonstrated that viral 5′ppp dsRNA structures (i.e. the panhandle) can bind and activate RIG-I even when encapsidated by nucleoprotein [21**]. Thus, apparently, RIG-I is capable of recognizing RNAs at all stages of infection. It is activated by the panhandle on viral nucleocapsids in the immediate early phase, that is directly after they entered the cytoplasm [21**], but also by newly synthesized viral RNAs later in infection [19,20].

Inhibition of RIG-I by segmented NSVs

For all the evolutionary advantages of having a segmented genome, these viruses face a particular problem: with every additional piece of RNA they increase their load of RIG-I ligands. The extreme is represented by influenza viruses, which expose no less than eight 5′ppp dsRNAs per infectious particle. Thus, strong and efficient anti-RIG-I measures are essential for these viruses to survive. The viral escape mechanisms act on many different levels (Figure 2). First of all, it might be no coincidence that influenza viruses and other orthomyxoviruses have evolved the unusual lifestyle of replicating in the nucleus. By this, they avoid constant exposure to cytoplasmic RIG-I, only passing the cytoplasm at the beginning and at the end of the replication cycle. Avoiding detection, in fact, represents an efficient way of RIG-I escape, since it is expected to demand less effort than blocking the amplified and diversified antiviral signalling events occurring further downstream. Influenza viruses exhibit also other features which could be interpreted as RIG-I escape. Firstly, the panhandle is not a perfect RIG-I ligand, but possesses mismatches and non-canonical base-pairs which weaken the dsRNA structure (see Figure 1c, left panel). Also, the nucleotide sequence of the orthomyxoviral panhandle enables the dsRNA panhandle to convert into a so-called ‘hook’ or ‘corkscrew’ by folding back the individual strands onto themselves [16,22] (see Figure 1c, right panel). The dsRNA-disrupting nucleotides and structural switches are known to regulate viral promoter activity, but it can be speculated they are disturbing RIG-I binding as well.

The panhandles of the less-segmented Bunyaviridae and Arenaviridae families often contain larger continuous double-stranded regions (see Figure 1d–f). However, some bunyavirus genera (Hantavirus and Nairovirus) are cleaving off the gamma and beta phosphates of the RIG-I-critical 5′ppp group [23] (see Figure 1e), and some arenavirus genome ends can exhibit a single 5′ overhanging nucleotide [24] with the potential to disturb RIG-I function [17] (see Figure 1f).

It must be noted, however, that most of the viral panhandle modifications are apparently insufficient for a complete innate immune escape. While genomic RNAs isolated from particles of those bunyaviruses which cleave off the 5′ppp do not activate RIG-I or antiviral signalling [23], the genomic RNAs of other bunyaviruses, influenza viruses, but also arenaviruses clearly activate RIG-I-dependent IFN induction if transfected into cells [23,25]. In the case of arenaviruses, this might be due to the fact that only a fraction of the RNAs actually seems to contain the unpaired 5′ nucleotide [24]. Moreover, RIG-I is most likely more flexible in its ligand requirements in vivo, a view which is supported by the observations that (i) not all nucleotides of the dsRNA helix are bound by amino acids of RIG-I [7,9**], (ii) bulges and 5′ overhangs of the 5′ppp dsRNA can be tolerated by RIG-I to some extent [10,13], and (iii) RNAs other than 5′ppp dsRNA can activate RIG-I [11,12,18*].

Independent of the 5′end, dsRNA from a certain length on is a prototypical hallmark of virus infection and a strong inducer of innate immune responses. In the cytoplasm, not only RIG-I but also the sister helicase MDA5 and the RNA-binding antiviral kinase PKR are activated by dsRNA [4,6]. NSVs avoid formation of dsRNA hybrids between the viral genome and its complementary RNA species (antigenome and mRNA) [14], most probably due to the separate packaging of genome and antigenome into nucleocapsids. Moreover, influenza viruses use the cellular helicases UAP56 and URIH49 to unwind any such dsRNA replication intermediates [26*].
RIG-I antagonism by segmented negative-strand RNA viruses. Infection with segmented NSVs generates structures with the potential to act as agonists of RIG-I. The viruses have therefore evolved escape strategies (shown in green) which act on all levels of RIG-I signalling (shown in light red). For details see text. Abbreviations: Gn: glycoprotein (N-terminal), IFN: type I interferon, IKKe: Inhibitor of nuclear factor kappa-B kinase subunit epsilon, IRF-3/7: interferon regulatory factors 3/7, ISG: interferon stimulated gene, MAVS: mitochondrial antiviral signalling protein, ML: matrix protein long, NP: nucleoprotein, NS1: non-structural protein 1, NSs: non-structural protein encoded on the S segment, NSV: negative-strand RNA virus, OTU: ovarian tumour domain, PA: polymerase acidic, PB: polymerase basic, RIG-I: retinoic acid-inducible gene I, SFTSV: severe fever with thrombocytopenia syndrome virus (a bunyavirus), TBK1: TANK-binding kinase 1, 5’ ppp dsRNA: 5’ triphosphorylated double-stranded RNA, TRIM: tripartite motif, Z: Zink finger protein. Dotted arrows symbolize signalling factors which have been left out for the sake of simplicity.
The above-described strategies of disrupting RIG-I-critical panhandle features and of preventing formation of large dsRNA stretches could be considered as basic, ‘passive’ abilities of segmented NSVs against the antiviral IFN system. On top of this, there are several active mechanisms in place. First of all, it is interesting to note that all segmented NSV families (but not the non-segmented ones) are performing ‘cap-snatching’, that is, they cleave capped oligonucleotides from host cell mRNAs and use them as primers for viral transcription. The responsible endonuclease domain is embedded in the viral polymerase and likely contributes to the downmodulation of antiviral cell responses [1]. In fact, it was recently discovered that influenza viruses can express the endonuclease domain alone, by ribosomal frameshift, and that this protein (PA-X) strongly suppresses host cell gene expression [27**]. The influenza virus polymerase subunits (PB1, PB2 and PA) can also block IFN upregulation independent of cap-snatching, by binding to MAVS [28,29]. Prime example of a separate influenza virus factor dedicated to IFN antagonism is the nonstructural protein NS1, a truly multifunctional protein known to bind dsRNA, RIG-I and its regulatory ubiquitin ligases TRIM 25 and Riplet, and PKR, but also to inhibit host cell mRNA processing, transport, and translation in a strain-specific manner [1,30**]. Some influenza A virus strains express an additional protein, PB1-F2, which also downmodulates innate immune reactions [31–33]. A related orthomyxovirus, Thogoto virus, has evolved a different strategy. It encodes a splice variant of the viral matrix protein, ML, which blocks IFN induction by associating with IRF-7 and the general transcription factor IIB [34,35]. The three-segmented Bunyaviruses similarly express IFN antagonists, most prominently the non-structural protein Ns1. Ns1 proteins of different bunyavirus genera either bind and degrade RIG-I [36], degrade PKR [37,38], sequester TBK1 [39], or — in most cases — suppress host cell mRNA synthesis by dysregulating, sequestering, or degrading key factors of RNA polymerase II transcription [15,40,41]. Moreover for some Han-taviruses it was shown that the glycoprotein Gn downregulates IFN induction by inhibiting the IRF-3 kinase TBK1 [42], and that the nucleo-protein inhibits the nuclear import of NF-κB [43]. Likewise, for the bunyavirus genus Nairovirus the ovarian tumour (OTU) domain of polymerase interferes with NF-κB activation by a de-ubiquitylation/de-ISGylation activity [44]. Also in the case of arenaviruses several IFN antagonists were described, first of all the nucleo-protein which degrades viral dsRNA [45,46], and inhibits NF-κB and the IRF-3 kinase IκKε [47,48]. Moreover, the Z protein of several arenaviruses directly binds and inhibits RIG-I [49].

Thus, the evolutionary advantage of dividing a viral RNA genome into several, genetically semi-autonomous pieces comes at some costs. It has to be counterbalanced by strong and efficient anti-RIG-I strategies acting on all possible levels. Nonetheless, at late stages of infection the viral counter mechanisms can be overrun by accumulating erroneous replication products, eventually tipping the balance towards activation of antiviral and inflammatory responses. In the extreme case, a cytokine storm develops causing the severe symptoms typical for so many of these viruses.

Conflicts of interest
No conflicts of interest declared.

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References and recommended reading
Papers of particular interest, published within the period of review, have been highlighted as:

| ● | of special interest |
| — | of outstanding interest |


5. References


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Appendix

I. Zusätzliche Publikationen


II. Kongressbeiträge

Mündliche Präsentationen

Human-adaptive influenza virus mutation PB2 627K contributes to evasion of nucleocapsid recognition by RIG-I

Activation of RIG-I by incoming RNA virus nucleocapsids containing a 5'-triphosphorylated genome

5. Europäischer Kongress für Virologie (ECV; European congress of Virology), Lyon/ Frankreich (2013)
Activation of RIG-I by incoming RNA virus nucleocapsids containing a 5'-triphosphorylated genome

Activation of RIG-I by incoming RNA virus nucleocapsids containing a 5’-triphosphorylated genome

Activation of RIG-I by incoming RNA virus nucleocapsids containing a 5’-triphosphorylated genome

Activation of RIG-I by incoming RNA virus nucleocapsids containing a 5’-triphosphorylated genome

Activation of RIG-I by incoming RNA virus nucleocapsids containing a 5’-triphosphorylated genome

**Posterpräsentation**

4. Internationales Influenza-Treffen, Münster (2014)
Human-adaptive influenza virus mutation PB2 627K contributes to evasion of nucleocapsid recognition by RIG-I

Protein expression profile of plasmeisin IV knock-out Plasmodium berghei parasite and immunoreactive proteins identification

Recruitment of plasma membrane beta-hexosaminidase in lipid rafts upon T-cell activation
III. Verzeichnis der akademischen Lehrer

IV. Danksagung


Meinen Freunden Anna, Anne, Silvana, Cathi, Johannes, Kathi, und Evelyn. Schon die Studienzeit war mit Eurer Unterstützung eine wunderbare Zeit. Trotz der weiten Entfernungen haben wir uns nicht aus den Augen verloren und Ihr wart und werdet für mich immer die Besten sein.


„Leider läßt sich eine wahrhafte Dankbarkeit mit Worten nicht ausdrücken.“

Johann Wolfgang von Goethe (1749 - 1832)